

Description

The invention relates to pharmaceutical compositions comprising a combination of a plasminogen activator and a thrombin inhibitor and to the use of said compositions for the prophylaxis or therapy of coagulation disorders.

Mammalian plasma contains two enzymatic systems coexisting in a dynamic equilibrium: the coagulation system which is capable of forming blood clots and the fibrinolytic system capable of dissolving blood clots. This dynamic equilibrium ensures that clots normally form only when and where they are required to prevent loss of blood from injured vessels and that after natural repair of the injury the superfluous clots are dissolved. In cases where the fibrinolytic capacity of the body is insufficient to remove intravascular clots (thrombi), for example in patients suffering from thromboembolisms or post-surgical complications, it may be indispensable to use exogenously administered fibrinolytic (thrombolytic) agents.

One component of the fibrinolytic system is a group of enzymes named plasminogen activators, which convert the zymogen plasminogen to the proteolytic enzyme plasmin. Plasmin then degrades the fibrin network of the clots to form soluble products.

Two types of plasminogen activators (hereinafter referred to as "PAs") can be isolated from human body fluids or cells: urokinase or urokinase-type plasminogen activator (hereinafter referred to as "u-PA"), a serine protease occurring e.g. in urine and kidney cells, and tissue-type plasminogen activator (hereinafter referred to as "t-PA") which is produced by endothelial cells and found in a number of endocrine tissues.

Both t-PA and u-PA exist in two molecular forms: a single-chain form (often designated as "sct-PA" and "scu-PA", respectively) and a two-chain (tc) form. The single-chain or pro-enzyme form is converted into the two-chain form by the action of proteolytic enzymes at well-defined positions in the polypeptide sequence. The resulting two chains of the processed PA protein remain attached to each other via a sulphur-sulphur bridge. The carboxyterminal fragment or B-chain mediates the enzymatic activity of the PA whereas the aminoterminal A-chain contains regulatory units such as fibrin binding sites.

While it is generally acknowledged that t-PA and u-PA (especially scu-PA) are of utmost usefulness in thrombolytic therapy it is nevertheless observed that none of these drugs alone is sufficient for permanently opening of occluded vessels. As the reocclusion rate is extraordinarily high additional therapeutical means are to be taken which exert a favourable influence on the course of disease.

Recently, thrombin inhibitors such as heparin, low molecular weight fractions of heparin or arginine derivatives (for examples argipidin) have been used in combination with plasminogen activators in anti-thrombotic therapy [cf. Y. Tamao et al., *Thromb. Haemost.* 56, 28-34 (1986); J.M. Stassen et al., *ibid.* 58, 947-950 (1987); European Patent Application No. 181,267]. The inhibition of thrombin has the following effects on the coagulation system: (1) the conversion of fibrinogen to monomeric fibrin is inhibited; (2) the maturation of factor XIII to factor XIIIa which converts soluble fibrin into the insoluble fibrin matrix, is prevented; soluble fibrin is more susceptible to proteolytic degradation by plasmin than insoluble fibrin; (3) the formation of factors Va and VIIIa (feed-back accelerators of thrombin formation) is inhibited; (4) plasminogen activator inhibitor 1 (PAI-1) is proteolytically degraded; while (5) thrombin bound in the thrombomodulin-thrombin complex is not inhibited by thrombin inhibitors; the complex can therefore continue to act as an activator of protein C which together with protein S proteolytically degrades factors Va and VIIIa (supra). The action of the plasminogen activator t-PA itself leads to local hyper-aggregability mediated by activated platelets [see R.J. Sheberski et al., *Thromb. Res.* 52, 381 (1988)]. Hence, thrombin inhibitors are believed to prevent the continuous formation of fibrin and the growth of thrombi by continuous deposition of newly formed fibrin. Moreover, since scu-PA is cleaved by thrombin at the Arg¹⁵⁶-Phe¹⁵⁷ bond to yield an inactive two chain form, it is inactive without concomitant administration of a thrombin inhibitor.

Also, during lysis significant amounts of fibrin are generated in circulation [Owen et al., *Blood* 72, 616-620 (1988); H.J. Rapold et al., *Circulation* 79, 980-988 (1988)]. Altogether, lysis via a plasminogen activator leads to an imbalance between thrombogenic and thrombolytic forces, locally as well as systemically, resulting in incomplete lysis of existing clots, rapid reocclusion at the same sites and novel clot formation at other predisposed sites. Anticoagulation by heparin will not prevent these side effects of lysis (Rapold, supra).

As indicated by the first results available the combined administration of plasminogen activator and thrombin inhibitor might help to overcome the reocclusion problem arising in conventional fibrinolytic plasminogen activator therapy. However, in view of the relatively small anticoagulant effects observed in the combined administration of heparin or arginine derivatives with plasminogen activators there is a need for a combination of plasminogen activator and a strong thrombin inhibitor having a more pronounced inhibitory effect on the coagulation cascade thus minimizing the reocclusion problem in fibrinolytic therapy. The combination with a strong thrombin inhibitor would also reduce the amount of the plasminogen activator

required to dissolve the clot and would further decrease the time required for dissolving the clot. This is especially important when life-threatening conditions are to be managed. It is an object of the invention to provide such an improved combination for use in a combined anticoagulant/fibrinolytic treatment of coagulation disorders.

It has surprisingly been found that the dissolution of thrombi is accelerated significantly and the risk of reocclusion is considerably reduced when the fibrinolytic plasminogen activator is applied in combination with hirudin which is known to be a well-tolerated and patent inhibitor of venous thrombosis and vascular occlusion [Markwardt et al., *Thromb. Haemost.* **47**, 226-229 (1982)].

Accordingly, the invention concerns a pharmaceutical combination composition comprising as component A a plasminogen activator and as component B hirudin together with a pharmaceutically acceptable carrier.

Plasminogen activators are human t-PA and human u-PA in the two-chain or, preferably, one-chain form including glycosylated, partially glycosylated or unglycosylated forms obtained from natural sources or via recombinant DNA techniques from transformed mammalian cells or transformed microorganisms, such as *Escherichia coli* or *Saccharomyces cerevisiae*. Examples of such plasminogen activators are those described by M. Winkler et al. in *Biochemistry* **25**, 4041-4045 (1986); B. Rotzkin et al., *Proc. Natl. Acad. Sci. USA* **78**, 3313-3317 (1981); D. Pennica et al., *Nature* **301**, 214-221 (1983); and in European Patent Applications No. 41,766, 92,182, 93,619 and 143,081.

The term "plasminogen activator" includes furthermore mutants of human u-PA and human t-PA, u-PA/t-PA hybrid plasminogen activators and mutants of the latter, especially single-chain mutants and hybrids, furthermore fragments thereof retaining the fibrinolytic activity.

Mutants are especially those mutants which render the PA protease resistant. Such mutants are covalently modified at sites of proteolysis by proteases occurring in blood such as thrombin or plasmin, so that they are no longer susceptible to protease hydrolysis at these locations. The target sites of u-PA include Lys¹³⁵ to Lys¹³⁶ (cleavage at this site generates the so-called low molecular weight form of scu-PA or "LUK"), Arg¹⁵⁶ to Phe¹⁵⁷ (susceptible to thrombin attack) and Lys¹⁵⁸ to Ile¹⁵⁹ (cleavage at this site by plasmin generates tzu-PA). Suitable u-PA mutants have site specific substitutions, insertions or deletions of amino acid residues at one or more of these target sites. Especially preferred are those mutants in which one amino acid residue or both amino acid residues forming the target sites are deleted or in which at least one of these amino acid residues is replaced by another amino acid residue so that the resulting mutants are resistant to proteolytic attack. For reference, see European Patent Applications No. 200,451, 210,279 and 288,435.

In further mutants of u-PA the unique N-glycosylation site occurring at Asn³⁰² (Asn³⁰²-Ser-Thr) is modified such that glycosylation cannot take place at this site.

Scu-PA and preferred scu-PA mutants have the formula I

Ser Asn Glu Leu His Gln Val

5 Pro Ser Asn Cys Asp Cys Leu Asn Gly Gly Thr Cys Val Ser Asn
 Lys Tyr Phe Ser Asn Ile His Trp Cys Asn Cys Pro Lys Lys Phe
 Gly Gly Gln His Cys Glu Ile Asp Lys Ser Lys Thr Cys Tyr Glu
 10 Gly Asn Gly His Phe Tyr Arg Gly Lys Ala Ser Thr Asp Thr Met
 Gly Arg Pro Cys Leu Pro Trp Asn Ser Ala Thr Val Leu Gln Gln
 Thr Tyr His Ala His Arg Ser Asp Ala Leu Gln Leu Gly Leu Gly
 Lys His Asn Tyr Cys Arg Asn Pro Asp Asn Arg Arg Arg Pro Trp
 15 Cys Tyr Val Gln Val Gly Leu Lys Pro Leu Val Gln Glu Cys Met
 Val His Asp Cys Ala Asp Gly X₁ X₂ Pro Ser Ser Pro Pro Glu
 Glu Leu Lys Phe Gln Cys Gly Gln Lys Thr Leu Arg Pro Y₁ Y₂
 20 Y₃ Ile Ile Gly Gly Glu Phe Thr Thr Ile Glu Asn Gln Pro Trp
 Phe Ala Ala Ile Tyr Arg Arg His Arg Gly Gly Ser Val Thr Tyr
 Val Cys Gly Gly Ser Leu Ile Ser Pro Cys Trp Val Ile Ser Ala
 Thr His Cys Phe Ile Asp Tyr Pro Lys Lys Glu Asp Tyr Ile Val
 25 Tyr Leu Gly Arg Ser Arg Leu Asn Ser Asn Thr Gln Gly Glu Met
 Lys Phe Glu Val Glu Asn Leu Ile Leu His Lys Asp Tyr Ser Ala
 Asp Thr Leu Ala His His Asn Asp Ile Ala Leu Leu Lys Ile Arg
 30 Ser Lys Glu Gly Arg Cys Ala Gln Pro Ser Arg Thr Ile Gln Thr
 Ile Cys Leu Pro Ser Met Tyr Asn Asp Pro Gln Phe Gly Thr Ser
 Cys Glu Ile Thr Gly Phe Gly Lys Glu Z₁ Ser Z₂ Asp Tyr Leu
 35 Tyr Pro Glu Gln Leu Lys Met Thr Val Val Lys Leu Ile Ser His
 Arg Glu Cys Gln Gln Pro His Tyr Tyr Gly Ser Glu Val Thr Thr
 Lys Met Leu Cys Ala Ala Asp Pro Gln Trp Lys Thr Asp Ser Cys
 40 Gln Gly Asp Ser Gly Gly Pro Leu Val Cys Ser Leu Gln Gly Arg
 Met Thr Leu Thr Gly Ile Val Ser Trp Gly Arg Gly Cys Ala Leu
 Lys Asp Lys Pro Gly Val Tyr Thr Arg Val Ser His Phe Leu Pro
 45 Trp Ile Arg Ser His Thr Lys Glu Glu Asn Gly Leu Ala Leu

in which X₁ and X₂ independently from each other represent Lys, an amino acid residue other than a basic amino acid residue or a chemical bond, Y₁ is Arg, an amino acid residue other than a basic amino acid residue or a chemical bond, Y₂ is Phe, an acidic amino acid residue or a chemical bond, Y₃ is Lys, an amino acid residue other than a basic amino acid residue or a chemical bond, Z₁ is Asn which is preferably yeast-specifically glycosylated, or is another amino acid residue, and Z₂ is Thr or another amino acid residue different from Ser.

The term "amino acid residue" is intended to embrace the residues of all genetically encoded amino acids, such as acidic amino acid residues, for example the residues of glutamic acid and aspartic acid, basic amino acid residues, for example the residues of arginine, lysine and histidine, and neutral amino acid residues, for example the residues of asparagine, glutamine, methionine, glycine, alanine, leucine, isoleucine, valine, serine, threonine, phenylalanine, tyrosine or proline.

In order to prevent glycosylation at the N-glycosylation site the tripeptide sequences recognised as signal for N-glycosylation -Asn-L-Thr(or Ser)- wherein Asn is the acceptor and L can be any of the 20 genetically encoded amino acids except proline or aspartic acid which impede glycosylation has to be altered. Replacement of the Asn (Z₁) and/or Thr (Z₂) residues in the above tripeptide sequence by any other amino acid would abolish formation of glycosidic linkages at this site. For convenience, modification of the N-glycosylation site is not done at the protein level. Instead, it is advantageous to modify the gene coding for scu-PA in such a way that upon expression of said modified gene by a host a mutant scu-PA is produced in which the N-glycosylation site is altered in such a way that glycosylation cannot take place at this site. For replacement those amino acids are preferred which have a similar size and polarity as the amino acids to be replaced. The number of possible substituents is limited by the given nucleotide sequence in the corresponding DNA and the possible codons. Especially, asparagine is replaced by valine, leucine, isoleucine, alanine, serine, threonine or, in particular, glutamine, and threonine by valine, methionine or, in particular, alanine. For reference, see European Patent Application No. 288,435.

Preferred scu-PA compounds of the formula I are those in which X₁ is Lys, Gly or Ser, X₂ represents Lys, Y₁ is Arg, Y₂ is Phe, Asp or Glu, Y₃ is Lys, Z₁ is Asn or Gln and Z₂ is Thr.

Corresponding preferred scu-PA mutants are [Gly¹³⁵]-scu-PA, [Ser¹³⁵]-scu-PA, [Asp¹⁵⁷]-scu-PA, [Ser¹³⁵,Asp¹⁵⁷]-scu-PA and [Gly¹³⁵,Asp¹⁵⁷]-scu-PA wherein Asn³⁰² is yeast specifically glycosylated, and furthermore [Gln³⁰²]-scu-PA, [Gly¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA and [Ser¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA.

The target site of t-PA with respect to proteolytic attack by proteases is in the sequence Phe²⁷⁴-Arg²⁷⁵-Ile²⁷⁶-Lys²⁷⁷. Suitable t-PA mutants have specific substitutions, insertions or deletions of amino acid residues at this site so that they are resistant to proteolytic attack (see above). For reference, see European Patent Applications No. 225,286, 227,462 and 233,013.

There are three sites for N-glycosidic linkage in the t-PA molecule: -Asn¹¹⁷-Ser-Ser-, Asn¹⁸⁴-Gly-Ser- and Asn⁴⁴⁸-Arg-Thr-.

T-PA and t-PA glycosylation mutants are, for example, those having the formula

Ser Tyr Gln Val Ile Cys Arg Asp Glu Lys Thr Gln Met Ile Tyr Gln Gln
His Gln Ser Trp Leu Arg Pro Val Leu Arg Ser Asn Arg Val Glu Tyr Cys
Trp Cys Asn Ser Gly Arg Ala Gln Cys His Ser Val Pro Val Lys Ser Cys
Ser Glu Pro Arg Cys Phe Asn Gly Gly Thr Cys Gln Gln Ala Leu Tyr Phe
Ser Asp Phe Val Cys Gln Cys Pro Glu Gly Phe Ala Gly Lys Cys Cys Glu
Ile Asp Thr Arg Ala Thr Cys Tyr Glu Asp Gln Gly Ile Ser Tyr Arg Gly
Thr Trp Ser Thr Ala Glu Ser Gly Ala Glu Cys Thr Asn Trp X₁ Ser Y₁
Ala Leu Ala Gln Lys Pro Tyr Ser Gly Arg Arg Pro Asp Ala Ile Arg Leu
Gly Leu Gly Asn His Asn Tyr Cys Arg Asn Pro Asp Arg Asp Ser Lys Pro
Trp Cys Tyr Val Phe Lys Ala Gly Lys Tyr Ser Ser Glu Phe Cys Ser Thr
Pro Ala Cys Ser Glu Gly Asn Ser Asp Cys Tyr Phe Gly X₂ Gly Y₂ Ala
Tyr Arg Gly Thr His Ser Leu Thr Glu Ser Gly Ala Ser Cys Leu Pro Trp
Asn Ser Met Ile Leu Ile Gly Lys Val Tyr Thr Ala Gln Asn Pro Ser Ala

Gln Ala Leu Gly Leu Gly Lys His Asn Tyr Cys Arg Asn Pro Asp Gly Asp
 Ala Lys Pro Trp Cys His Val Leu Lys Asn Arg Arg Leu Thr Trp Glu Tyr
 5 Cys Asp Val Pro Ser Cys Ser Thr Cys Gly Leu Arg Gln Tyr Ser Gln Pro
 Gln Phe Arg Ile Lys Gly Gly Leu Phe Ala Asp Ile Ala Ser His Pro Trp
 Gln Ala Ala Ile Phe Ala Lys His Arg Arg Ser Pro Gly Glu Arg Phe Leu
 Cys Gly Gly Ile Leu Ile Ser Ser Cys Trp Ile Leu Ser Ala Ala His Cys
 10 Phe Gln Glu Arg Phe Pro Pro His His Leu Thr Val Ile Leu Gly Arg Thr
 Tyr Arg Val Val Pro Gly Glu Glu Glu Gln Lys Phe Glu Val Glu Lys Tyr
 Ile Val His Lys Glu Phe Asp Asp Asp Thr Tyr Asp Asn Asp Ile Ala Leu
 15 Leu Gln Leu Lys Ser Asp Ser Ser Arg Cys Ala Gln Glu Ser Ser Val Val
 Arg Thr Val Cys Leu Pro Pro Ala Asp Leu Gln Leu Pro Asp Trp Thr Glu
 Cys Glu Leu Ser Gly Tyr Gly Lys His Glu Ala Leu Ser Pro Phe Tyr Ser
 20 Glu Arg Leu Lys Glu Ala His Val Arg Leu Tyr Pro Ser Ser Arg Cys Thr
 Ser Gln His Leu Leu X₃ Arg Y₃ Val Thr Asp Asn Met Leu Cys Ala Gly
 Asp Thr Arg Ser Gly Gly Pro Gln Ala Asn Leu His Asp Ala Cys Gln Gly
 Asp Ser Gly Gly Pro Leu Val Cys Leu Asn Asp Gly Arg Met Thr Leu Val
 25 Gly Ile Ile Ser Trp Gly Leu Gly Cys Gly Gln Lys Asp Val Pro Gly Val
 Tyr Thr Lys Val Thr Asn Tyr Leu Asp Trp Ile Arg Asp Asn Met Arg Pro

(II)

in which X₁, X₂ and X₃ each represent Asn or another genetically encoded amino acid, Y₁ and Y₂ each
 represent Ser or another genetically encoded amino acid other than Thr and Y₃ represents Thr or another
 35 genetically encoded amino acid other than Ser.

More particularly, the t-PA proteins have the formula II in which X₁, X₂ and X₃ each represent Asn and
 Y₁, Y₂ and Y₃ have the meanings given above.

Genetically encoded amino acids are those specified above.

For replacement of Asn in position X₁, X₂ and/or X₃ amino acids Gln, Thr and Ser are preferred. For
 40 replacement of Ser Y₁ and/or Y₂ and for the replacement of Thr Y₃ amino acids Ala and Asn are preferred.

Especially, only one of the radicals X₁ and Y₁ and/or only one of the radicals X₂ and Y₂ and/or only one
 of the radicals X₃ and Y₃ is replaced by another amino acid.

In particular, the t-PA mutants have the formula II in which one, two or three of the radicals X₁, X₂ and
 X₃ represent Asn and the other(s) represent(s) an amino acid residue selected from Gln, Thr and Ser, Y₁
 45 and Y₂ each represent Ser and Y₃ represents Thr, or in which X₁, X₂ and X₃ each represent Asn, one or
 two of the radicals Y₁ and Y₂ represent Ser and the other(s) is (are) selected from the group consisting of
 Ala and Asn and Y₃ represents Thr, or in which X₁, X₂ and X₃ each represent Asn, Y₁ and Y₂ each
 represent Ser and Y₃ is selected from the group consisting of Ala and Asn.

Especially, the t-PA mutants have the formula II in which X₁, X₂ and X₃ each represent Asn, and at
 50 least one of radicals Y₁, Y₂ and Y₃ is Ala or Asn and the other(s) is (are) Ser (Y₁ and Y₂) or Thr (Y₃).

Examples of such t-PA mutants are [Asn¹¹⁹]-t-PA, [Ala¹⁸⁶]-t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹, Ala¹⁸⁶]-t-PA,
 [Asn¹¹⁹, Ala¹⁸⁶, Asn⁴⁵⁰]-t-PA and [Asn¹¹⁹, Ala¹⁸⁶, Ala⁴⁵⁰]-t-PA.

Fragments of u-PA and t-PA are especially those in which segments or domains of natural u-PA or t-PA
 which are not essential for fibrinolytic activity are deleted. Such fragments and processes for their
 55 preparation are known per se.

U-PA/t-PA hybrid plasminogen activators are especially single-chain hybrid PAs comprising an amino
 acid sequence containing all or discrete A-chain domains of human u-PA or discrete A-chain domains of
 human u-PA and human t-PA, linked in series to the catalytic region of human t-PA and single-chain hybrid

PAs comprising an amino acid sequence containing all or discrete A-chain domains of human t-PA or discrete A-chain domains of human t-PA and u-PA, linked in series to the catalytic region of human u-PA as disclosed in European Patent Application No. 277,313 or a single-chain hybrid PA differing from u-PA in that the amino acid segment following the u-PA kringle up to the u-PA activation site is replaced by the amino acid segment following the t-PA kringle 2 up to the t-PA activation site.

Preferred hybrid PAs contain the catalytic region of human u-PA.

In particular, the single-chain hybrid PAs comprise an amino acid sequence selected from the group consisting of an amino acid sequence containing all of the A-chain domains of human t-PA, an amino acid sequence containing discrete A-chain domains of human t-PA, such as the finger domain or a kringle, especially the kringle 2, domain of human t-PA, and an amino acid sequence containing two, three or four A-chain domains of human t-PA and/or human u-PA, especially two or three domains of human t-PA or two or three domains of human u-PA and human t-PA, such as the finger, growth factor and kringle 2 domains of human t-PA, the finger and kringle 2 domains of human t-PA or the u-PA growth factor and t-PA kringle 2 domains, which amino acid sequence is linked in series to the catalytic region of human u-PA, and an amino acid sequence containing the u-PA growth factor and t-PA kringle 2 domains which amino acid sequence is linked in series to the catalytic region of human t-PA.

Preferably, the hybrid PA amino acid sequence starts with the N-terminal amino acids 1 to 5 of t-PA or the N-terminal amino acids 1 to 12 of u-PA or starts with any junction sequence naturally N-terminally linked to the first domain of the hybrid PA or with a fragment of such a junction sequence which fragment preferably has at least five amino acid residues.

In the hybrid PAs the A-chain domains are connected via natural junction sequences, fused junction sequences or hybrid junction sequences or fragments thereof. Thus, a first domain is linked to a second domain by the junction sequence naturally occurring at the C-terminus of the first domain, by the junction sequence naturally occurring at the N-terminus of the second domain, by a fused junction sequence composed of said junction sequences or by fragments thereof.

The A-chain domains of the hybrid PAs according to the invention are linked to the B-chain serine protease region by a junction sequence selected from the group consisting of the junction sequence linking the kringle 2 domain to the B-chain in human t-PA, the junction sequence linking the kringle domain to the B-chain in human u-PA and a hybrid sequence composed of subsequences of said junction sequences wherein said junction sequence includes a processing site capable of being cleaved by plasmin and, N-terminally thereto, a cysteine residue which can participate in a sulphur-sulphur bridge to the catalytic B-chain region, the junction sequence preferably having at least fourty and up to 60 amino acid residues.

Most preferred is the junction of the domains at a position which is defined by the exon-intron junctions on the corresponding DNA. The junction of the A-chain to the B-chain is most preferably at the activation site.

In particular, the single chain hybrid plasminogen activators are selected from the group consisting of such a hybrid plasminogen activator comprising the A-chain of u-PA or an A-chain essentially consisting of the u-PA growth factor and the t-PA kringle 2 domains linked in series to the catalytic region (B-chain) of t-PA, and a hybrid plasminogen activator comprising the A-chain of t-PA, an A-chain essentially consisting of the finger domain of t-PA, an A-chain essentially consisting of the u-PA growth factor and t-PA kringle 2 domains, an A-chain essentially consisting of the t-PA finger and kringle 2 domains or an A-chain essentially consisting of the t-PA finger, growth factor and kringle 2 domains said A-chain being linked in series to the catalytic region (B-chain) of u-PA, wherein the A-chain is linked to the B-chain via a junction sequence comprising an activation site and a cysteine residue capable of forming a sulphur-sulphur bond to the B-chain or the single chain hybrid plasminogen activator comprises an A-chain essentially consisting of the t-PA kringle 2 domain linked to the catalytic region (B-chain) of u-PA at the activation site.

Especially preferred is a single chain hybrid plasminogen activator selected from the group consisting of such a hybrid PA comprising an A-chain essentially consisting of the u-PA growth factor domain and the t-PA kringle 2 domain linked in series to the catalytic region (B-chain) of t-PA, and a hybrid PA comprising an A-chain essentially consisting of the t-PA kringle 2 domain or of the t-PA finger and kringle 2 domains linked in series to the catalytic region (B-chain) of u-PA, wherein the junction between the A-chain domain(s) and the B-chain is at the activation site.

Preferred hybrid PAs according to the invention are

[uPA(1-158)-tPA(276-527)],

[uPA(1-131)-tPA(263-527)],

[tPA(1-275)-uPA(159-411)],

[tPA(1-262)-uPA(132-411)],

[uPA(1-44)-tPA(176-261)-uPA(134-411)],

[tPA(1-49)-tPA(262-275)-uPA(159-411)],
 [tPA(1-49)-uPA(134-411)],
 [tPA(1-49)-tPA(176-275)-uPA(159-411)],
 [tPA(1-49)-tPA(176-262)-uPA(132-411)],
 5 [uPA(1-44)-tPA(176-527)],
 [uPA(1-44)-tPA(176-275)-uPA(159-411)],
 [uPA(1-133)-tPA(262-275)-uPA(159-411)],
 [tPA(1-3)-tPA(176-275)-uPA(159-411)],
 [tPA(1-86)-tPA(176-275)-uPA(159-411)] and
 10 [tPA(1-86)-tPA(176-262)-uPA(132-411)].

Especially preferred are hybrid plasminogen activators [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] and [tPA(1-3)-tPA(176-275)-uPA(159-411)].

In the mutants of the hybrid PAs at least one, preferably all, of the N-glycosylation sites are modified such that glycosylation cannot take place at this (these) site(s).

15 Especially, asparagine is replaced by valine, leucine, isoleucine, alanine or, in particular, glutamine, and serine or threonine by valine, methionine or, in particular, alanine.

Preferred modified hybrid PAs are

[tPA(1-49)-tPA(262-275)-uPA(159-301, Gln, 303-411)],
 [tPA(1-49)-tPA(176-185, Ala, 187-275)-uPA(159-301, Gln, 303-411)],
 20 [uPA(1-44)-tPA(176-185, Ala, 187-449, Ala, 451-527)],
 [tPA(1-3)-tPA(176-185, Ala, 187-275)-uPA(159-301, Gln, 303-411)],
 [tPA(1-86)-tPA(176-185, Ala, 187-275)-uPA(159-301, Gln, 303-411)],
 [tPA(1-49)-tPA(176-275)-uPA(159-301, Gln, 303-411)],
 [tPA(1-3)-tPA(176-275)-uPA(159-301, Gln, 303-411)],
 25 [uPA(1-44)-tPA(176-449, Ala, 451-527)],
 [tPA(1-86)-tPA(176-275)-uPA(159-301, Gln, 303-411)],
 [uPA(1-133)-tPA(262-275)-uPA(159-301, Gln, 303-411)],
 [uPA(1-133)-tPA(262-275)-uPA(159-303, Asn, 305-411)],
 [uPA(144)-tPA(176-185, Ala, 187-275)-uPA(159-301, Gln, 303-411)] and
 30 [uPA(1-44)-tPA(176-185, Ala, 187-275)-uPA(159-303, Asn, 305-411)].

The term "hirudin" includes hirudins obtained from natural sources, such as from the leech *Hirudo medicinalis*, hirudins obtained via recombinant DNA techniques, as well as mutants and fragments thereof retaining the unique thrombin inhibitory activity. A number of such hirudins have been published by D. Tripiet [Folia Haematol. 115, 30 (1988)].

35 The hirudin component of the compositions according to the invention is, for example, hirudin variant HV1 [D. Bagdy et al., Methods Enzymol. 45, 669-678 (1976); US Patent No. 4,745,177] having the formula

Val Val Tyr Thr Asp Cys Thr Glu Ser Gly
 40 Gln Asn Leu Cys Leu Cys Glu Gly Ser Asn
 Val Cys Gly Gln Gly Asn Lys Cys Ile Leu
 Gly Ser Asp Gly Glu Lys Asn Gln Cys Val
 Thr Gly Glu Gly Thr Pro Lys Pro Gln Ser
 45 His Asn Asp Gly Asp Phe Glu Glu Ile Pro
 Glu Glu ^WTyr Leu Gln

(III)

50 wherein W represents the phenolic hydroxy group of Tyr (desulfato hirudin HV1) or a group of the formula -O-SO₃H (hirudin HV1), hirudin variant HV2 [cf. R.P. Harvey et al., Proc. Natl. Acad. Sci. USA 83, 1084-1088 (1986)] having the formula

Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 5 Ile Leu Gly Ser Asn Gly Lys Gly Asn Gln Cys Val Thr Gly
 Glu Gly Thr Pro Asn Pro Glu Ser His Asn Asn Gly Asp Phe
 Glu Glu Ile Pro Glu Glu Tyr Leu Gln

(IV)

and hirudin variant PA [cf. J. Dodt et al., Biol. Chem. Hoppe-Seyler 367, 803-811 (1986)] having the formula

Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 15 Ile Leu Gly Ser Gln Gly Lys Asp Asn Gln Cys Val Thr Gly
 20 Glu Gly Thr Pro Lys Pro Gln Ser His Asn Gln Gly Asp Phe
 Glu Pro Ile Pro Glu Asp Ala Tyr Asp Glu

(V)

or a fragment or a mutant of any of said hirudin variants.

Especially preferred is desulfato hirudin HV1.

30 Suitable fragments of such hirudin variants, especially of hirudin variant HV1, are known, for example from FEBS Lett. 211, 10 (1987), J. Med. Chem. 31, 1009 (1988), Biochemistry 27, 8170 (1988), J. Biol. Chem. 264, 8692 (1989) and from EP 276014, and consist, for example, of the 9 to 30 C-terminal amino acids of hirudin HV1 or PA. Further fragments include, for example, hirudin HV1 lacking the C-terminal amino acid Gln or the C-terminal dipeptide residue Leu-Gln.

35 Mutants of the hirudin according to the invention include hirudins, especially hirudin HV1, carrying site specific mutations at the N-terminus, in the core region or at the C-terminus. Such N-terminal mutants of hirudin HV1 are, for example, [Ile^{1,2}]-desulfato hirudin HV1 (in which the N-terminal amino acids Val-Val are replaced by Ile-Ile) or [Gly¹]-desulfato hirudin HV1 (in which the N-terminal amino acid Val is replaced by Gly). Corresponding C-terminal mutants of hirudin HV1 include, for example, [Gln^{57,58,61,62}]-desulfato hirudin
 40 HV1 (in which amino acids 57, 58, 61 and 62 have been replaced by Gln) and [Pro⁶⁶]-desulfato hirudin HV1 (carrying an additional Pro residue at the C-terminus). Corresponding core mutants of desulfato hirudin HV1 include, for example, [Gln²⁷,Arg⁴⁷]-desulfato hirudin HV1 which additionally may carry an extra Met residue at the N-terminus ([Met⁰,Gln²⁷,Arg⁴⁷]-desulfato hirudin HV1) and [Gln²⁷,Gln³⁶,Arg⁴⁷]-desulfato hirudin HV1. A further mutant is [des-Val¹,Thr²]-desulfato hirudin HV1. A mutant of hirudin HV2 is, for example, [Lys⁴⁷]-
 45 hirudin HV2. All of these mutants are known or can be prepared by methods known in the art, for example by subjecting the gene coding for the respective parent hirudin to site-specific mutagenesis and expressing the mutated hirudin gene in a suitable host such as yeast.

Further mutants of said hirudins are, for example, those disclosed in European Patent Application No. 273,800.

50 In particular, the invention concerns a pharmaceutical composition comprising as component A a plasminogen activator selected from the group consisting of

(1) scu-PA,

(2) t-PA,

(3) a sct-PA mutant of the formula II in which X₁, X₂ and X₃ each represent Asn, and at least one of radicals Y₁, Y₂ and Y₃ is Ala or Asn and the other(s) is (are) Ser (Y₁ and Y₂) or Thr (Y₃), and

55 (4) a single chain u-PA/t-PA hybrid plasminogen activator selected from such a hybrid PA comprising an A-chain essentially consisting of the u-PA growth factor domain and the t-PA kringle 2 domain linked in series to the catalytic region (B-chain) of t-PA, and a hybrid PA comprising an A-chain essentially

consisting of the t-PA kringle 2 domain or of the t-PA finger and kringle 2 domains linked in series to the catalytic region (B-chain) of u-PA, wherein the junction between the A-chain domain(s) and the B-chain is at the activation site,

and as component B desulfato hirudin HV1, together with a pharmaceutically acceptable carrier.

5 In most preferred pharmaceutical compositions according to the invention the component A is selected from the group consisting of scu-PA, t-PA, [Ala450]-t-PA, [Asn119, Ala186, Ala450]-t-PA, [uPA(1-44)-tPA-(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] and [tPA(1-3)-tPA(176-275)-uPA(159-411)], and component B is desulfato hirudin HV1.

10 The plasminogen activators and hirudins according to the invention are known compounds. Especially preferred are those PAs and hirudins which have been prepared by recombinant DNA techniques making use of transformed mammalian or yeast cells, as described, for example, in European Patent Applications No. 143,081, 225,286, 200,655, 225,633 and 277,313.

15 The novel combination compositions according to the invention can be used in mammals (humans or animals) for the prevention or treatment of thrombosis or diseases caused by thrombosis, arteriosclerosis, myocardial and cerebral infarction, venous thrombosis, thromboembolism, post-surgical thrombosis, thrombophlebitis, etc.

The pharmaceutical compositions according to the invention can be used for the treatment of the above-mentioned indications when they are administered parenterally, such as intravenously or, as regards hirudin, also subcutaneously.

20 There are suitable infusion or injection solutions, preferably aqueous isotonic solutions or suspensions, it being possible to prepare these before use, for example from lyophilised preparations that contain the active ingredient(s) alone or together with a pharmaceutically acceptable carrier, such as mannitol, lactose, dextrose, human serum albumin and the like. The pharmaceutical compositions are sterilized and, if desired, mixed with adjuncts, for example preservatives, stabilisers, emulsifiers, solubilisers, buffers and/or salts (such as 0.9 % sodium chloride) for adjusting the isotonicity. Sterilization can be achieved by sterile
25 filtration through filters of small pore size (0.45 μ m diameter or smaller) after which the composition can be lyophilised, if desired. Antibiotics may also be added in order to assist in preserving sterility. For example, standard formulation techniques developed for the administration of PAs (i.e. European Patent Applications No. 93,619, 41,766, 112,940 and others) or novel compositions conferring sufficient solubility to the PA in
30 low pH buffered media (examples are in European Patent Application No. 211,592, German Offenlegungsschrift No. 3,617,752, 3,617,753, 3,642,960) may be used. Hirudins are as such highly soluble in aqueous solutions and need no additives for enhanced solubility. For stability reasons it is recommended to reconstitute lyophilized samples in 5 % dextrose or 5 % mannitol or 0.9 % sodium chloride and the like.

35 The combination compositions according to the invention are in a state to allow the active ingredients (plasminogen activator and hirudin) to be administered (e.g. infused) at the same time and by the same route (i.e. cannula) or to apply first hirudin, preferably by bolus injection, and then, starting within 5 to 10 minutes thereafter, the second component, especially the plasminogen activator, preferably by infusion, or, in particular, to apply first the total dose of hirudin together with a small quantity, for example 5 to 20 %, of the plasminogen activator by bolus injection and then, starting within 5 to 10 minutes thereafter, the main
40 quantity of the plasminogen activator by infusion. Infusion is done within 30 minutes to 3 hours.

Depending upon the type of the disease and the age and the condition of the patient, the daily dose to be administered once for the treatment of a patient weighing approximately 70 kg is in the range from 10 to 200, especially 20 to 100 mg, plasminogen activator and 5 to 100 mg, especially 10 to 30 mg, hirudin [especially a dosis which leads to a times two prolongation of activated partial thromboplastin time (aPTT)].
45 Accordingly, the weight ratio between the plasminogen activator and the hirudin in the composition may vary, in general between 20:1 and 2:1. A weight ratio between 5:1 and 2:1 is preferably used.

The pharmaceutical compositions according to the present invention are dispensed in unit dosage forms, for example in ampoules comprising therapeutically effective amounts of both components (plasminogen activator and hirudin) or, preferably, in double ampoules comprising therapeutically effective
50 amounts of the components separately, together with a pharmaceutically acceptable carrier.

The present pharmaceutical compositions are produced in a manner known per se applying conventional lyophilising or dissolving procedures, for example, if necessary, by mixing the plasminogen activator and the hirudin and optionally the pharmaceutically acceptable carrier, and, for the preparation of a lyophilisate, freeze-drying an aqueous solution obtained. The compositions contain from approximately 0.1
55 % to 20 %, especially from approximately 1 % to 10 %, and in the case of lyophilisates up to 100 % of the active ingredients.

The invention relates also to the use of the combination compositions according to the invention for the prophylactic and therapeutic treatment of the human or animal body, especially for the above-mentioned

clinical syndromes, in particular for the prophylaxis and therapy of thrombosis or diseases caused by thrombosis in the human or animal body.

A further object of the present invention is founded on the surprising observation that hirudin is a powerful agent which dramatically reduces the reocclusion (new formation of thrombi) rate of vessels (veins or arteries) which have previously been opened by mechanical (PTCA) or chemical (lysis) means and thus improves the prospects of definitely curing the coagulation disorder. Accordingly, the invention concerns a method for preventing the reocclusion of previously occluded and mechanically or chemically reopened blood vessels in a mammal comprising administering to said mammal a therapeutically effective amount of hirudin. Preferably, the administration of hirudin follows immediately after the opening of the occluded vessels, for example after termination of the thrombolytic therapy using, for example, plasminogen activators optionally in combination with antithrombotic agents. Depending upon the type of the disease and the age and the condition of the patient, the daily dose of hirudin to be administered by infusion or, preferably, by two bolus injections, for the treatment of a patient weighing approximately 70 kg is in the range from 20 to 100 mg, especially about 50 to 70 mg. The anti-reocclusion therapy using hirudin is continued until the risk of reocclusion has minimized, for example for about 1 to 3 weeks, in particular for about 2 weeks.

Brief description of the drawings

In the following experimental part various embodiments of the present invention are described with reference to the accompanying drawings in which:

Fig. 1 shows the fibrin accretion during dissolution of the clot by t-PA in the presence of heparin and desulfato hirudin, respectively.

Fig. 2 shows the fibrin accretion during dissolution of the clot by t-PA in the presence of various concentrations of desulfato hirudin.

Fig. 3 schematically illustrates the construction of plasmid pML350.

Fig. 4 schematically shows the plasmid map of plasmid pJDB207/GAPFL-HIR.

The following Examples illustrate the invention without implying any limitation.

Example 1: In vitro clot lysis

Human blood is directly withdrawn into 3.8 % sodium citrate solution (ratio blood/citrate solution 9:1). The mixture is centrifuged at 1,000 rpm for 10 min to yield citrated plasma.

A mixture consisting of

150 µl citrated plasma

125 µl 0.9 % sodium chloride solution

100 µl desulfato hirudin HV1 or MCI-9038 (see below) in 0.9 % sodium chloride

is prepared directly in 1 ml cuvettes equilibrated at 37 °C. The mixture is incubated for 15 min at 37 °C. Then, 25 µl of a 1 µg/ml t-PA stock solution (American Diagnostica, New York) and 300 µl 0.0025 M calcium chloride solution are added to start coagulation.

The change in the optical density (coagulation) is continuously measured at 608 nm in a Uvikon 810 P spectrophotometer equipped with a kinetic program.

The time (in minutes) from addition of the CaCl₂ solution to a change in absorbance greater than 0.025 is defined as the coagulation time. The clot lysis time (in minutes) is defined as the time from the beginning of the clotting to the time at which the absorbance decreased to 10 % of the maximum absorbance.

The thrombin inhibitor MCI-9038 [(2R,4R)-4-methyl-1-[N²-(3-methyl-1,2,3,4-tetrahydro-8-quinolyl)-sulfonyl]-L-arginyl]-2-piperidine-carboxylic acid] is chemically synthesized as described in S. Okamoto et al., Biochem. Biophys. Res. Commun. 101, 440-446 (1981). It is used at the following end concentrations: 0.07, 0.14, 0.28, 0.56 and 1.1 µM. The desulfato hirudin used has been prepared by recombinant DNA technology using transformed yeast according to the procedure described in European Patent Application No. 225,633. It is used at the following end concentrations: 0.8, 1.6, 3.2, 8 and 16 nM. The concentrations chosen of MCI-9038 and desulfato hirudin are equipotent concentrations in terms of prolongation of clotting time (coagulation), e.g. 1.2 µM MCI-9038 and 16 nM desulfato hirudin lead to the same prolongation of aPTT.

It is evident that in order to achieve the same effect as exhibited by a given concentration in desulfato hirudin, the twenty-fold concentration in MCI-9038 has to be taken.

Example 2: Rabbit jugular vein thrombosis model

The experiments are performed with New Zealand white rabbits (weighing 2.3-3.5 kg) anaesthetized by intravenous (marginal ear vein) injection of 30 mg/kg sodium pentobarbital.

5 Through a paramedical incision of the neck, the jugular veins and the right carotid artery are exposed. The right carotid artery is cannulated with a 1.5 mm diameter polyethylene cannula for blood sampling. The two jugular veins are isolated, cleared over a distance of 2 cm, small side branches are ligated. Each jugular segment is emptied of blood and blood flow is temporarily occluded both proximally and distally by two angiostaths, 1.5 cm apart. A 10 cm length 3.0 TiCron braided polyester suture, presoaked in a collagen
10 solution, is introduced lengthwise in the lumen of the isolated jugular vein for a distance of 4 cm to avoid the embolization of the thrombus to be produced. 150 µl of citrated rabbit blood containing 5 µl of ¹²⁵I-fibrinogen (approx. 500 000 cpm) is then aspirated in a 1 ml tubercoline syringe containing 50 µl (5 I.U.) of thrombin and 10 µl of CaCl₂ (0. 25M). The components are quickly mixed and injected into the isolated segment of the jugular vein via a 25 gauge needle. Thrombi are allowed to age for 30 min and then both
15 vessel clamps are removed and blood flow restored.

a) Enhancement of t-PA induced lysis by desulfato hirudin HV1

Intravenous infusions are carried out by using a constant rate infusion pump. t-PA derived from melanoma cell cultures is obtained from BioResponse, USA. A total dose of 0.5 mg t-PA is infused through
20 the marginal ear vein over a period of 3 hours. Along with t-PA either saline, standard heparin (KabiVitrum, Sweden, 140 USP/mg; 0.75 mg/kg) or hirudin (1 mg/kg) are infused. Doses of heparin and hirudin are used which give in control experiments the same prolongation in aPTT (activated partial thromboplastin time), e.g. times two.

25 As a control a group of animals is only infused with saline without addition of t-PA. At the end of the infusion, the thrombi remaining in the vessel are removed and their size is determined by measuring the amount of residual ¹²⁵I-labeled fibrinogen remaining in the thrombi and comparing the results with those of the saline controls.

The results of the experiment are summarized in Table 1.

Table 1:

35	Combination	% lysis	Mean of n animals
	First set of experiments		
40	t-PA + saline	44	n = 5
	t-PA + desulfato hirudin	52	n = 12

45	Second set of experiments		
	t-PA + saline	37	n = 18
50	t-PA + heparin	34	n = 18

b) Inhibition of thrombus accretion by desulfato hirudin HV1 in t-PA treated rabbits

55 The effects of standard heparin and recombinant desulfato hirudin in preventing thrombus extension during thrombolysis with t-PA are assessed as their ability to inhibit the accretion of ¹²⁵I-fibrinogen onto autologous non-radioactive thrombi preformed in the jugular vein of rabbits. Standard sized non-radioactive

thrombi are produced as described above. 15 min after the thrombi are formed each animal is injected with 10 μ Ci of 125 I-fibrinogen. Five min later the infusion of t-PA along with standard heparin or desulfato hirudin is started (concentrations as in Example 2a). At the end of the three hours infusion, the venous segments containing the thrombi are tied off, split open longitudinally and the remaining thrombi are removed. The specific activity of the whole blood fibrinogen is estimated from the mean of the blood samples collected at hourly intervals throughout the infusion. The ratio of the radioactivity of the thrombus to the circulating fibrinogen radioactivity is used to quantify thrombus size, which is then expressed as μ g or 125 I-fibrinogen accreted onto the thrombus.

The results are summarized in Figure 1 and Figure 2.

Figure 1 shows that even during the dissolution of the clot new 125 I-fibrinogen $\rightarrow$ fibrin is deposited on the surface of the preformed clot (e.g. 52 μ g with only t-PA). Heparin has clearly no beneficial effect (49 versus 52 μ g, statistically not significant), whereas desulfato hirudin shows a clear reduction in thrombus accretion (SH: standard heparin; S.D.: standard deviation; r-HIR: recombinant desulfato hirudin).

Figure 2 shows that with increasing concentrations of desulfato hirudin this effect can even be potentiated.

The same effects are observed when a plasminogen activator different from t-PA, such as scu-PA or a u-PA/t-PA hybrid plasminogen activator, or another hirudin compound, such as desulfato hirudin PA or a hirudin mutant, is used in the combination treatment. Corresponding hirudin mutants can be prepared as described in Example 3.

Example 3: Production of desulfato hirudin HV1 mutants

A. Construction of the plasmid pML350 (see Fig. 3)

a) Digestion of the DNA of plasmid pIN-III-ompA-2

10 μ g of plasmid pIN-III-ompA-2 [J. Ghayeb et al., EMBO-J. 3, 2437 (1984)] are dissolved in 25 μ l of 100 mM Tris-HCl pH 7.5, 50 mM NaCl and 100 μ g/ml gelatine and are digested with the restriction endonucleases EcoRI and BamHI. The solution is adjusted to TNE and extracted with phenol/chloroform. The DNA is precipitated with ethanol. The vector DNA pIN-III-ompA-2/EcoRI/BamHI is isolated after electrophoresis in agarose by gel elution.

b) Digestion of the DNA of plasmid pML310

20 μ g of the plasmid pML310 (see European Patent Application No. 168342) are digested in 50 μ l of 100 mM Tris-HCl pH 7.5, 50 mM NaCl and 100 μ g/ml gelatine with the restriction endonucleases EcoRI and BamHI. The solution is adjusted to TNE and extracted with phenol/chloroform. The DNA is precipitated with ethanol. The F₁-F₂-DNA (hirudin gene) is isolated after gel electrophoresis in agarose by gel elution.

c) Ligation of the F₁-F₂-DNA (hirudin gene) from pML310 with the vector DNA pIN-III-ompA-2/EcoRI/BamHI

1 μ g of F₁-F₂-DNA (hirudin gene)/EcoRI/BamHI and 30 μ g of the vector DNA pIN-III-ompA-2/EcoRI/BamHI are dissolved in 50 μ l of 100 mM Tris-HCl pH 7.5, 50 mM NaCl and 100 μ g/ml gelatine, adjusted to TNE. The solution is extracted with phenol/chloroform and the DNA is precipitated with ethanol. The DNA precipitate is dissolved in 20 μ l of a solution of 50 mM Tris-HCl (pH 7.8), 10 mM MgCl₂, 10 mM DTT, 0.5 mM ATP, and 100 μ g/l gelatine and treated with 25 units/ μ l T₄ DNA ligase (Biolabs) at 15 °C for 3 h. By this way the recombinant plasmid pML350 is created, which contains the F₁-F₂-DNA (hirudin gene) inserted.

d) Transformation of E. coli HB101 with plasmid pML350

The E. coli HB101 cells pretreated with calcium that are prepared as described by Mandel et al. [J. Mol. Biol. 53, 159 (1970)].

The solution obtained in c), which contains the recombinant plasmid pML350, is heated at 65 °C for 10 min in order to inactivate the T₄ DNA ligase and is then cooled to 37 °C. 10 μ l of the resulting reaction mixture are added to 150 μ l of calcium-treated E. coli HB101 cells in 10 mM MgCl₂ and 10 mM Tris-HCl (pH 7.5) in a total volume of 200 μ l.

Subsequently, this mixture is cooled in ice for 30 min, heated for 2 min at 42 °C and then left to stand for 50 min in 1 ml of L-medium (Bacto tryptone 10 g/l; Bacto yeast extract 5 g/l; NaCl 5 g/l; glucose 5 g/l; ampicillin 0.1 g/l) at 37 °C. The mixture is then spread out in aliquots of 0.2 ml on 5 agar plates (McConkey Agar, Difco), which contain 60 µg/ml of ampicillin (Serva). The agar plates are then maintained at 37 °C for 16-18 hours. 185 ampicillin resistant colonies of transformed E. coli HB101 cells are obtained.

e) Screening the colonies that contain F₁-F₂-DNA

6 transformed colonies are pressed off onto nitrocellulose filter B85 (Schleicher and Schull). In accordance with Grunstein and Hogness [Proc. Natl. Acad. Sci. USA 72, 3961 (1979)] the colonies are lysed and their denatured DNA is fixed on the filter. Subsequently prehybridization of the filters is carried out in 20 ml (per filter) of 4xSET [= solution of 30 mM Tris-HCl (pH 8), 150 mM NaCl, 1 mM EDTA], 0.1 % (w/v) Ficoll 400 (Pharmacia), 0.5 % SDS, 50 µg/ml denatured calf thymus DNA for 4 h at 64 °C. Subsequently the nitrocellulose filters are treated in 20 ml (per filter) of 5xSET (w/v), 0.1 % (w/v) Ficoll 400, 0.2 % SDS and 50 µl/ml denatured calf thymus DNA for 16 h at 64 °C with the ³²P radioactively labelled probe (approximately 10³-10⁴ Cerencov cpm per filter). A mixture consisting of oligonucleotides 46/64 complementary, 1/58, 96/67 and 154/64 complementary (see European Patent Application No. 168342) is used as probe.

Subsequently, the filters are washed twice in 2xSET, 0.2 % SDS at room temperature, then twice in 2xSET, 0.5 % SDS at 60 °C (first for 30 min, then for 60 min). The filters are then dried between 3 MM paper (Whatman) and placed at -80 °C on an X-ray film (Fuji) with an intensifying screen (Ilford) for 1-2 days.

The resulting autoradiogram shows 5 positive colonies (clones) which can be used for further processing, one of which received the designation pML350.

B. Construction of plasmid pBH109

As plasmid pML350 originates from plasmid pIN-III-ompA-2 including a multi-cloning linker (EcoRI, HindIII, BamHI sites) it contains 12 additional base pairs in front of the mature hirudin gene coding for Ala, Gln, Phe, Met. To get mature desulphatohirudin expressed these 12 base pairs are looped out by in vitro mutagenesis making use of a 27mer oligonucleotide.

a) Preparation of pML350/XbaI/BamHI (SI)

5 µg of plasmid pML350 are digested with the endonucleases XbaI and BamHI. The larger of the two XbaI-BamHI fragments (SI) is isolated by gel elution after electrophoresis on agarose and dissolved in 1 mM Tris-HCl pH 7.5, 0.1 mM EDTA.

b) Preparation of pML350/PvuI (SII)

5 µg of plasmid pML350 are digested with the endonuclease PvuI. The linearized DNA pML350/PvuI is subsequently digested with 3 units of intestinal alkaline phosphatase (Boehringer) for 30 min at 37 °C. The enzyme is inactivated by heating the solution for 60 min at 65 °C. The linear pML350/PvuI (SII) DNA is isolated by gel elution after electrophoresis on agarose and dissolved in 1 mM Tris-HCl pH 7.5, 0.1 mM EDTA.

c) Preparation of the oligonucleotide (27 mer) I27

In analogy to the procedure described in European Patent Application No. 168342 the following DNA fragment (designated I27) has been synthesized:

5'-GTA GCG CAG GCC GTT GTT TAC ACC GAC-3'

I27

The phosphorylation at the 5' ends was done with [γ-³²P]-ATP and T₄ polynucleotide kinase (Boehringer) as described [Molecular Cloning, A Laboratory Manual (ed. T. Maniatis et al.), Cold Spring Harbor Lab. (1982),

p. 125].

d) Construction of plasmid pBH109

5 0.3 µg each of SI DNA and of SII DNA are mixed with 40 pmol of the phosphorylated DNA fragment 127 in 27 µl of 1 mM Tris-HCl pH 7.5, 0.1 mM EDTA. To the mixture 3 µl of 10X polymerase-ligase buffer (1M NaCl, 65 mM Tris-HCl pH 7.5, 80 mM MgCl₂ and 10 mM β-mercaptoethanol) are added. This mixture is heated for 3 min in a boiling water bath to denature the DNA fragments. Subsequently, the mixture is gradually cooled (about 1 °C/min) to 30 °C and incubated at this temperature for 30 min. Further the mixture is incubated at 4 °C for 30 min and afterwards for 10 min in ice.

10 12 µl of the four deoxyribonucleotide phosphates (7.5 mM each), 6 µl of 10 mM ATP, 6 µl of T₄ DNA ligase (2.5 U/µl) and 1.2 µl of Klenow DNA polymerase (Boehringer, 5 U/µl) are added and the DNA mixture (total volume 55 µl) is incubated for 16 h at 12.5 °C.

15 The DNA mixture is digested with 2 units of endonuclease EcoRI for 1 h at 37 °C in order to destroy unchanged starting plasmid pML350. With this procedure, plasmid pBH109 is formed. Plasmid pBH109 contains the lpp promoter and the lac promoter/operator operably linked to the ompA-2 signal sequence linked in frame to the gene coding for mature desulphato-hirudin.

e) Transformation of E. coli HB101 with plasmid pBH109

20 The transformation with calcium treated E. coli HB101 cells is done as described above. The total reaction mixture used is 55 µl.

f) Screening of the colonies which contain plasmid pBH109

25 100 transformed colonies are cultivated, from each colony plasmid DNA is prepared and digested with EcoRI. All plasmid DNAs, which are not digestable with EcoRI are potent plasmids pBH109 which is lacking the EcoRI site.

Two positive identical colonies have been identified. One of them is selected and designated pBH109.

30 The correct sequence of the F₁-F₂-DNA following the ompA-2 leader sequence is confirmed by sequence analysis.

C. Mutation of the residue Lys27 of hirudin to Asn27 using single stranded M13mp19/hirudin.

35

	24	26	28	30
coding strand	Gln	Gly	Asn	<u>Lys</u> Cys Ile Leu
of hirudin	5'CAG	GGT	AAC	<u>AAA</u> TGC ATC CTG 3'

40

mutagenic

primer 1	3'GTC	CCA	TTG	<u>TTA</u> ACG TAG GAC 5'
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45

mutated	5'CAG	GGT	AAC	<u>AAT</u> TGC ATC CTG 3'
coding strand	Gln	Gly	Asn	<u>Asn</u> Cys Ile Leu

50 Mutagenic primers are synthesised using the phosphoramidite method [M.H. Caruthers, in Chemical and Enzymatic Synthesis of Gene Fragments (H.G. Gassen and A. Lang, Eds.) Verlag Chemie, Weinheim, Federal Republic of Germany] on an Applied Biosystems (Model 380B) synthesiser.

55

I. Preparation of M13mp19/hirudin.XbaI-BamHI cut M13mp19 DNA.

5 To 5 μ l M13mp19 double stranded DNA (ds-DNA; 0.1 μ g/ml; BRL) are added 2 μ l React 2 (500 mM Tris-HCl, pH 8.0; 100 mM $MgCl_2$, 500 mM NaCl) (BRL), 1 μ l XbaI (10 U/ μ l), 0.5 μ l BamHI (10 U/ μ l) and 12 μ l H_2O . After incubation at 37°C for 1.5 h, 0.5 μ l BamHI, 2.5 μ l React 3 (500 mM Tris-HCl, pH 8.0; 100 mM $MgCl_2$; 1000 mM NaCl) (BRL), and 2 μ l H_2O are added and incubation is continued at 37°C for 1 h. The volume is made up to 100 μ l with H_2O . The ds-DNA is isolated by phenol extraction and ethanol precipitation, and dissolved in 30 μ l of TE buffer (Tris-HCl 10 mM, EDTA 1 mM, pH 8.0).

Insert DNA.

15 Five μ g of the plasmid pBH109 are cut with XbaI and BamHI as described above and the digest is electrophoresed for 3 h at 150 volt using a 3.5 % polyacrylamide gel with 1x TBE buffer (10x TBE buffer: 108 g Tris, 55 g boric acid, 9.3 g EDTA $\cdot$ 2 H_2O /l). The XbaI-BamHI fragment containing the hirudin gene (250 bp) is visualised under UV light after immersing the gel in 400 ml 1x TBE buffer containing 10 μ l of ethidium bromide solution (10 μ g/ml in water). The part of the gel containing the restriction fragment is cut from the gel and placed in a dialysis bag with 500 μ l of 0.5x TBE, and the DNA is electroeluted in a BIO-
20 RAD minigel electrophoresis apparatus using 0.5x TBE as the running buffer at 170 volt for 30 min. The DNA is loaded onto an Elutip-d column (Schleicher & Schull) equilibrated with 0.5x TBE. The column is washed with 2 ml of 0.5x TBE and the DNA is eluted with 1 M NaCl in 0.5x TBE (1 ml). The DNA is precipitated with ethanol and redissolved in 10 μ l of TE buffer.

25 Ligation of XbaI-BamHI hirudin insert into M13mp19 and preparation of single stranded DNA.

Five μ l XbaI-BamHI hirudin insert, 2 μ l XbaI-BamHI cut M13mp19, 1 μ l 10x ligase buffer (50 mM Tris-HCl, pH 7.5, 10 mM $MgCl_2$, 10 mM dithio-threitol), 1 μ l ATP, 1.5 μ l T4 DNA ligase (BRL; 1 U/ μ l) are mixed and incubated overnight at 14°C. Five μ l of ligation mixture are used to transform *E. coli* JM101 competent cells according to the method of J. Messing [Methods in Enzymology 101, 21-78 (1983)]. Twelve clear
30 plaques are picked and single stranded DNA (ss-DNA) is prepared from each plaque as described by J. Messing (supra). The DNA designated M13mp19/hirudin is redissolved in 50 μ l of TE buffer (0.1-0.5 μ g/ μ l).

II. Site-directed mutagenesis.35 Phosphorylation of mutagenic primer.

200 pmol (23 μ l) of mutagenic primer 1 (see above) is phosphorylated by adding 3 μ l 10x kinase buffer (1M Tris-HCl, 0.1M $MgCl_2$, 0.1M dithio-threitol, pH 8.3) 3 μ l 10 mM ATP and 1 μ l T4 polynucleotide kinase (BRL, 10 U/ μ l). After incubation at 37°C for 1 h, the reaction is stopped by heating at 65°C for 10 min.

Annealing of the mutagenic primer 1 to the single-stranded M13mp19/hirudin template.

45 Six μ l (0.5 μ g) of single-stranded M13mp19/hirudin is incubated with 3 μ l (20 pmol) of the phosphorylated mutagenic oligodeoxyribonucleotide (6.6 pmol/ μ l) and 1 μ l buffer A (0.2M Tris-HCl, pH 7.5, 0.1M $MgCl_2$, 0.5M NaCl, 0.01M DIT) at 70°C for 5 min, and cooled slowly to room temperature over 30 min.

Extension-ligation reaction.

50 To the above annealed mix is added 1 μ l buffer B (0.2M Tris-HCl, pH 7.5, 0.1M $MgCl_2$, 0.01M DTT), 1 μ l 10mM ATP, 4 μ l 2mM dNTP mixture, 5 μ l T4 DNA polymerase (Boehringer, 1 U/ μ l), 5 μ l T4 DNA ligase (BRL, 1 U/ μ l). This mixture is incubated at 16°C for 3 h. The reaction is stopped by incubating at 65°C for 10 min.

Transformation and preparation of single-stranded mutant DNA

The ligation mixture is diluted 1:20 with sterile H₂O, and 1 µl, 5 µl, as well as 1 µl undiluted ligation mixture is used to transform competent *E. coli* BMH 71-81 mut S cells [B. Kramer, W. Kramer and H.-J. Fritz, Cell 38, 879-887 (1984)]. The cells are plated out as described in the "M13 cloning and sequencing Handbook" (published by Amersham). Twelve colourless plaques are picked and ss-DNA is prepared as described above.

Screening of single-stranded DNA for mutant.

To screen for mutated single-stranded DNA, each of the 12 ss-DNA samples is sequenced by the dideoxynucleotide chain termination method [F. Sanger, S. Nickler and A.R. Coulson, Proc. Natl. Acad. Sci. USA 74, 5463-5467 (1977)]. Initially, only the dideoxynucleotide complementary to the expected mutated base is used in the reaction. Subsequently, the ss-DNA from several positive mutants are sequenced to establish the full DNA sequence of the mutant using T7 DNA polymerase (Sequenase, USB) following the method of Tabor and Richardson [Proc. Natl. Acad. Sci. USA 84, 4767-4771 (1987)]. The expected base change encoding Lys → Asn mutation at position 27 of the recombinant hirudin are observed in the DNA sequence. The phage DNA having the expected sequence is designated M13mp19/hirudin K27N.

Preparation of replicative form (RF) DNA from single-stranded M13mp19/hirudin K27N phage DNA.

Competent *E. coli* JM101 cells are transformed with 10-20 ng of single-stranded hirudin K27N mutant DNA and ds-DNA is prepared as described in the "M13 cloning and sequencing Handbook" (published by Amersham). A yield of 40-50 µg of ds-DNA is obtained from 100 ml of culture.

Isolation of mutant hirudin XbaI-BamHI insert.

The mutated hirudin XbaI-BamHI insert is cut out of 25 µg of the ds-DNA and purified as described in section IB. The DNA is dissolved in 20 µl of sterile water.

Preparation of XbaI-BamHI cut pIN-III-ompA-2 vector DNA

A digest of approximately 1.5 µg pIN-III-ompA-2 plasmid is made by adding 6 µl React 2 buffer (BRL), 2 µl (20 Units) XbaI, 1 µl BamHI (10 Units), 1 µl EcoRI (10 Units) and 37 µl H₂O (total volume 50 µl), and incubation for 3 h at 37°C. 1 µl (10 Units) BamHI, 1 µl (10 Units) EcoRI, 5 µl React 3 (BRL) and 12 µl H₂O are added and incubation continued for 1 h at 37°C.

Ligation of mutant hirudin K27N XbaI-BamHI insert DNA into XbaI-BamHI cut pIN-III-ompA-2 plasmid.

Nine µl hirudin K27N XbaI-BamHI insert DNA, 2 µl XbaI-BamHI cut pIN-III-ompA-2 vector DNA, 3 µl 10x ligation buffer (BRL) and 1 µl (1 U/µl) T4 DNA ligase (BRL) are mixed and incubated at 14°C for 16-20 h.

III. Expression of mutant [Asn²⁷]-desulfato hirudin in *E. coli* JM101. Transfection into *E. coli* strain JM101.

Five µl of ligation mixture is used to transform 0.3 ml of *E. coli* JM101 competent cells according to the method of J. Messing (supra). Three ml of 2xYT/Ampicillin (50 µg ampicillin/ml 2xYT) is added to the sample and the cells allowed to grow at room temperature for 1 h. A 1 ml sample of the culture is then taken and poured onto an LB/Ampicillin (50 µg ampicillin/ml LB-agar) plate and grown overnight at 37°C. The transforming plasmid DNA is referred to as pIN-III-ompA-2/HIR-K27N.

Selection of [Asn²⁷]-desulfato hirudin expressing colonies.

Ten bacterial colonies from the LB/Ampicillin plates are picked and grown separately for 5 h at 37°C in 5ml LB/Ampicillin (50 µg Ampicillin/ml LB). 1 ml samples are taken from each culture tube and the cells recovered by centrifugation (3000xg for 5 min). Each sample of cells is osmotically shocked by treatment with 100 µl of 10 mM Tris-HCl, pH 8.1 for 30 min at 0°C to release the material in the periplasmic space of the bacteria. The cells are removed by centrifugation as before, and the supernatant is tested for hirudin

activity. The sample which gives the highest inhibitory activity is selected for batch culture.

Batch culture and isolation of [Asn²⁷]-desulfato hirudin.

5 The remaining quantity (4 ml) of cells from the most active sample is used to inoculate 1 l of LB/Ampicillin (50 µg Ampicillin/ml LB). The culture is grown overnight at 37 °C, and the cells are recovered by centrifugation (3000xg for 15 min.). The cell pellet is osmotically shocked by resuspension in 50 ml of 10 mM Tris-HCl, pH 8.1 at 0 °C for 1 h. The cells are removed from the periplasmic fraction by centrifugation at 6000xg for 10 min.

10 Purification of [Asn²⁷]-desulfato hirudin.

15 The pH of the periplasmic fraction is adjusted to 6.5 with 0.1M HCl and filtered through a 0.45 µm filter (Nalgene). The protein is loaded onto a Mono-Q column FPLC system (Fast Protein Liquid Chromatography, Pharmacia-LKB) equilibrated with 50mM bis-Tris-HCl buffer pH 6.5. The desulfato hirudin mutant is eluted from the column using a linear salt gradient of 0-300 mM NaCl in bis-Tris-HCl pH 6.5 over 45 min. 0.8 ml fractions of the column eluate are collected and tested for hirudin activity as described above. The desulfato hirudin mutant-containing fractions are pooled, filtered as above and chromatographed on a Millipore-Waters HPLC system using a Brownlee Labs C8 reversed-phase HPLC column equilibrated with 20 0.09 % (v/v) trifluoroacetic acid in H₂O. The hirudin mutant is eluted from the column with a linear gradient of 7 to 28 % (v/v) acetonitrile in 0.09 % (v/v) trifluoroacetic acid in H₂O. [Asn²⁷]-desulfato hirudin having a purity of about or more than 98 % elutes as a single peak at 28 min.

D. Mutation of the residue Lys 27 of hirudin to Gln 27, and of the residue Lys 47 to Arg 47A: Mutation of Lys 27 to Gln 27

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27

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coding strand Gln Gly Asn Lys Cys Ile Leu
 of hirudin 5' CAG GGT AAC AAA TGC ATC CTG 3'

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mutagenic 3' GTC CCA TTG GTT ACG TAG GAC 5'
 primer 2A

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mutated 5' CAG GGT AAC CAA TGC ATC CTG 3'
 coding strand Gln Gly Asn Gln Cys Ile Leu

B: Mutation of Lys 47 to Arg 47

25

47

30

coding strand Gly Thr Pro Lys Pro Gln Ser
 of hirudin 5' GGT ACC CCG AAA CCG CAG TCT 3'

mutagenic 3' CCA TGG GGC TCT GGC GTC AGA 5'
 primer 2B

35

mutated 5' GGT ACC CCG AGA CCG CAG TCT 3'
 coding strand Gly Thr Pro Arg Pro Gln Ser

40 The procedures given above are repeated using mutagenic primers 2A and B to obtain and characterize the desired mutant protein in which Lys 27 is replaced by Gln 27 and Lys 47 by Arg 47. The transforming plasmid DNA is referred to as pIN-III-ompA-2/HIR-K27Q,K47R. The mutant is designated [Gln²⁷,Arg⁴⁷]-desulfato hirudin.

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E. Mutation of the residue Lys 27 of hirudin to Gln 27, the residue Lys 36 to Gln 36, and of the residue Lys 47 to Arg 47.

a: Mutation of Lys 36 to Gln 36

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		34	36	38	
	coding strand	Asp Gly Glu	<u>Lys</u>	Asn Gln Cys	
10	of hirudin	5' GAC GGT GAA	<u>AAA</u>	AAC CAG TGC	3'
	mutagenic				
15	primer 3	3' CTG CCA CTT	<u>GTT</u>	TTG GTC ACG	5'
	mutated				
	5'	GAC GGT GAA	<u>CAA</u>	AAC CAG TGC	3'
20	coding strand	Asp Gly Glu	<u>Gln</u>	Asn Gln Cys	

b: Mutation of Lys 27 to Gln 27

25 The mutation of Lys 27 to Gln 27 is performed according to D (above).

c: Mutation of Lys 47 to Arg 47

The mutation of Lys 47 to Arg 47 is performed according to D (above).

30 The procedures given above are repeated using mutagenic primers 2A, 3 and 2B to obtain and characterize the desired mutant protein in which Lys 27 is replaced by Gln 27, Lys 36 is replaced by Gln 36 and Lys 47 is replaced by Arg 47. The transforming plasmid DNA is referred to as pIN-III-ompA-2/HIR-K27Q,K36Q,K47R. The mutant is designated [Gln²⁷,Gln³⁶,Arg⁴⁷]-desulfato hirudin.

35 F. Extension of the N-terminus of [Gln²⁷,Arg⁴⁷]-desulfato hirudin with a methionine residue

		1	3	5	
	coding strand	signal seq.	Val Val Tyr Thr Asp Cys		
40	of hirudin	5' GCG CAG GCC ...	GTT GTT TAC ACC GAC TGC	3'	
	mutagenic				
45	primer 4	3' CGC GTC CGG	<u>TAC</u>	CAA CAA ATG TGG CTG ACG	5'
	mutated				
	5'	GCG CAG GCC	<u>ATG</u>	GTT GTT TAC ACC GAC TGC	3'
50	coding strand	signal seq.	<u>Met</u>	Val Val Tyr Thr Asp Cys	

The procedures given above are repeated using mutagenic primers 2A, 2B and 4 to obtain and characterize the desired mutant protein in which the N-terminus of [Gln²⁷,Arg⁴⁷]-desulfato hirudin is extended by Met. The protein is designated methionyl-[Gln²⁷,Arg⁴⁷]-desulfato hirudin.

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G. Mutation of the residues Glu 57,58,61,62 of hirudin to Gln 57,58,61,62

			56	58	60	62	64	
5	coding strand	Gly Asp Phe	<u>Glu</u>	<u>Glu</u>	Ile Pro	<u>Glu</u>	<u>Glu</u>	Tyr Leu Gln
	of hirudin	5' GTT GAC TTC	<u>GAA</u>	<u>GAA</u>	ATC CCG	<u>GAA</u>	<u>GAA</u>	TAC CTG CAG 3'
10	mutagenic							
	primer 5	3' CCA CTG AAG	<u>GTT</u>	<u>GTT</u>	TAG GGC	<u>GTT</u>	<u>GTT</u>	ATG GAC GTC 5'
15	mutated	5' GGT GAC TTC	<u>CAA</u>	<u>CAA</u>	ATC CCG	<u>CAA</u>	<u>CAA</u>	TAC CTG CAG 3'
	coding strand	Gly Gly Phe	<u>Gln</u>	<u>Gln</u>	Ile Pro	<u>Gln</u>	<u>Gln</u>	Tyr Leu Gln

The procedures given above are repeated using mutagenic primer 5 to obtain and characterize the desired mutant protein in which Glu 57,58,61,62 are replaced by Gln 57,58,61,62. The mutant is designated [Gln^{57,58,61,62}]-desulfato hirudin.

H. Construction of a yeast expression plasmid coding for [Pro⁶⁶]-desulfato hirudin

The DNA sequence coding for desulfato hirudin is extended by the oligonucleotide CCA, which codes for proline. The resulting new desulfato hirudin is expressed in yeast. It contains 66 aminoacids with a proline at its C-terminal end. This polypeptide is referred to as [Pro⁶⁶]-desulfato hirudin.

Yeast expression plasmid pJDB207/GAPFL-HIR (see Fig. 4; European patent application No. 225633) is digested with restriction endonucleases Sall and EcoRI. The 478 bp Sall-EcoRI fragment is separated on a 0.8 % preparative agarose gel in TBE buffer (90 mM Tris-base, 90 mM boric acid, 2.5 mM EDTA pH 8.3). The ethidiumbromide-stained fragment is isolated from the gel. The DNA is electroeluted in 0.2xTBE buffer for 45 min at 100 mA and purified by DE52 (Whatman) ion exchange chromatography. The DNA is eluted from the DE52 column with a high salt buffer (1.5 M NaCl, 10 mM Tris-HCl pH 8.0, 1 mM EDTA), precipitated with ethanol and redissolved in H₂O at a concentration of 0.1 pmoles/μl. The 478 bp Sall-EcoRI fragment contains the Sall-BamHI sequence of pBR322 fused to the BglII-EcoRI GAPFL promoter fragment.

Plasmid pJDB207/GAPFL-HIR is digested with BamHI and Sall. The large, 6.7 kb vector fragment is isolated as described above. The small, 740 bp Sall-BamHI fragment is also isolated. It contains the sequence of the 478 bp Sall-EcoRI fragment (see above) in addition to the PH05 signal sequence fused in frame to the coding sequence of desulfato hirudin. The 740 bp fragment is digested with AsuII. The DNA is extracted with phenol/chloroform, precipitated with ethanol and redissolved in H₂O.

A synthetic oligodeoxynucleotide of the formula

Pro

(1) 5'-CGAAGAAATCCCGGAAGAATACCTGCAGCCATAG -3'

(2) 3'- TTCTTTAGGGCCTTCTTATGGACGTCGGTATCCTAG -5'

is kinased in 40 μl of 60 mM Tris-HCl pH 7.5, 10 mM MgCl₂, 5 mM DTT, 0.5 mM ATP and 27 U of T4 polynucleotide kinase (Boehringer) for 45 min at 37°C. The reaction mixtures for oligonucleotides (1) and (2) are combined. The mixture is heated for 10 min at 75°C and allowed to cool to room temperature. The annealed oligonucleotide linker (1+2) is stored at -20°C.

0.85 μg (3.8 pmoles) of the AsuII digested DNA are incubated for 16 h at 15°C with a 100 fold excess of the kinased and annealed oligonucleotide linker over DNA ends in 150 μl of 60 mM Tris-HCl pH 7.5, 10 mM MgCl₂, 5 mM DTT, 3.5 mM ATP and 1200 U of T4 DNA ligase (Biolabs). After inactivation of the T4 DNA ligase for 10 min at 85°C the excess linkers are removed by precipitation of the DNA in the presence of 10 mM EDTA, 300 mM sodium acetate pH 6.0 and 0.54 volumes of isopropanol. The DNA is digested with EcoRI and BamHI. The resulting fragments are separated on a 2 % preparative agarose gel in TBE

buffer. The 262 bp fragment is recovered from the gel by electroelution and ethanol precipitation. The DNA is resuspended at a concentration of 0.1 pmoles/ μ l. The EcoRI-BamHI fragment contains the coding sequence of desulfato hirudin with the additional CCA triplet coding for Pro 66.

Three DNA fragments, isolated as described above, are ligated in the following reaction: 0.2 pmoles of the 478 bp Sall-EcoRI fragment, 0.2 pmoles of the 262 bp EcoRI-BamHI fragment and 0.1 pmoles of the 6.7 kb vector fragment are ligated in 10 μ l of 60 mM Tris-HCl pH 7.5, 10 mM MgCl₂, 5 mM DTT, 1 mM ATP and 200 U of T4 DNA ligase for 6 h at 15°C. A one μ l aliquot of the ligation mixture is added to 100 μ l of calcium-treated, transformation competent *E. coli* HB101 cells.

12 transformed, ampicillin-resistant colonies are grown in LB medium containing 100 mg/l of ampicillin. Plasmid DNA is prepared [Holmes et al., Anal. Biochem. 114 (1981), 193] and analysed by BamHI, EcoRI double digests. The presence of the mutation in the DNA is confirmed by DNA sequencing [Sanger et al., Proc. Natl. Acad. Sci. USA 74 (1977) 5463]. One clone with the CCA codon in the hirudin structural gene is referred to as pJDB207/GAPFL-HIR (Pro 66).

1. Mutation of the residues Val 1,2 of hirudin to Ile 1,2

			1		3		5	
coding strand	signal seq.	<u>Val</u>	<u>Val</u>	Tyr	Thr	Asp	Cys	
of hirudin	5'	GCG	CAG	GCC	<u>GTT</u>	<u>GTT</u>	TAC	ACC GAC TGC 3'
mutagenic								
primer 6	3'	CGC	GTC	CGG	<u>TAA</u>	<u>TAA</u>	ATG TGG CTG	ACG 5'
mutated	5'	GCG	CAG	GCC	<u>ATT</u>	<u>ATT</u>	TAC	ACC GAC TGC 3'
coding strand	signal seq.	<u>Ile</u>	<u>Ile</u>	Tyr	Thr	Asp	Cys	

The procedures given above are repeated using mutagenic primer 6 to obtain and characterize the desired mutant protein in which the N-terminal amino acids Val 1,2 are replaced by Ile 1,2. The mutant is designated [Ile^{1,2}]-desulfato hirudin.

J. Mutation of the residues Val 1 of hirudin to Gly 1

			1		3		5	
coding strand	signal seq.	<u>Val</u>	Val	Tyr	Thr	Asp	Cys	
of hirudin	5'	GCG	CAG	GCC	<u>GTT</u>	GTT	TAC	ACC GAC TGC 3'
mutagenic								
primer 7	3'	CGC	GTC	CGG	<u>CCA</u>	CAA	ATG TGG CTG	ACG 5'
mutated	5'	GCG	CAG	GCC	<u>GGT</u>	GTT	TAC	ACC GAC TGC 3'
coding strand	signal seq.	<u>Gly</u>	Val	Tyr	Thr	Asp	Cys	

The procedures given above are repeated using mutagenic primer 7 to obtain and characterize the desired mutant protein in which the N-terminal amino acid Val 1 is replaced by Gly 1. The mutant is designated [Gly¹]-desulfato hirudin.

In an analogous manner applying the procedures detailed in Example 3 any other desired mutant of desulfato hirudin HV1, hirudin HV2 and hirudin PA can be produced.

Example 4: Pharmaceutical composition for parenteral administration and application

The pharmaceutical composition consists of seven vials each containing 10 mg lyophilised activator t-PA and 300 mg dextrose and an eighth vial containing 30 mg of lyophilised desulfato hirudin HV1 and 100 mg dextrose.

A solution for parenteral administration of t-PA is prepared by dissolving the contents of each of the seven vials in one 10 ml ampoule of sterile pyrogene-free solvent containing 100 mM ammonium acetate buffer, adjusted to pH 4 with acetic acid.

A solution for parenteral administration of desulfato hirudin is prepared by dissolving the contents of the vials in 3 ml of sterile 0.9 % sodium chloride solution.

10 % of the total plasminogen activator dosis (i.e. 7 mg t-PA) in 7 ml buffer and 30 mg desulfato hirudin in 3 ml buffer are aspirated into a sterile syringe containing 10 ml sterile 0.9 % sodium chloride solution and intravenously given as an initial loading dose over 5 min. The remaining 63 mg of t-PA are injected into a 250 ml bottle of sterile 0.9 % sodium chloride solution and then intravenously infused over 90 min to achieve opening of the occluded vessel.

Alternatively, 7 mg of t-PA (7 ml) and 10 mg of desulfato hirudin (1 ml) dissolved as described (supra) are given as a bolus dosis over 5 min. The remaining 63 mg of t-PA and 20 mg of desulfato hirudin are infused over 90 min in a total volume of 315 ml 0.9 % sodium chloride solution.

Instead of t-PA it is also possible to use the same amount of another plasminogen activator mentioned hereinbefore, such as scu-PA or a u-PA/t-PA hybrid plasminogen activator. Analogously, instead of desulfato hirudin HV1 it is also possible to use the same amount of another hirudin mentioned hereinbefore, such as hirudin PA or a mutant of desulfato hirudin HV1.

Claims

1. A pharmaceutical composition comprising in separate unit dosage forms as component A a plasminogen activator selected from the group consisting of u-PA and t-PA including glycosylated, partially glycosylated or unglycosylated forms thereof, a u-PA/t-PA hybrid plasminogen activator and mutants and fragments thereof retaining the fibrinolytic activity and as component B hirudin together with pharmaceutically acceptable carriers.
2. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator in the single chain form.
3. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator of the formula



In an analogous manner applying the procedures detailed in Example 3 any other desired mutant of desulfato hirudin HV1, hirudin HV2 and hirudin PA can be produced.

Example 4: Pharmaceutical composition for parenteral administration and application

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The pharmaceutical composition consists of seven vials each containing 10 mg lyophilised activator t-PA and 300 mg dextrose and an eighth vial containing 30 mg of lyophilised desulfato hirudin HV1 and 100 mg dextrose.

10 A solution for parenteral administration of t-PA is prepared by dissolving the contents of each of the seven vials in one 10 ml ampoule of sterile pyrogene-free solvent containing 100 mM ammonium acetate buffer, adjusted to pH 4 with acetic acid.

A solution for parenteral administration of desulfato hirudin is prepared by dissolving the contents of the vials in 3 ml of sterile 0.9 % sodium chloride solution.

15 10 % of the total plasminogen activator dosis (i.e. 7 mg t-PA) in 7 ml buffer and 30 mg desulfato hirudin in 3 ml buffer are aspirated into a sterile syringe containing 10 ml sterile 0.9 % sodium chloride solution and intravenously given as an initial loading dose over 5 min. The remaining 63 mg of t-PA are injected into a 250 ml bottle of sterile 0.9 % sodium chloride solution and then intravenously infused over 90 min to achieve opening of the occluded vessel.

20 Alternatively, 7 mg of t-PA (7 ml) and 10 mg of desulfato hirudin (1 ml) dissolved as described (supra) are given as a bolus dosis over 5 min. The remaining 63 mg of t-PA and 20 mg of desulfato hirudin are infused over 90 min in a total volume of 315 ml 0.9 % sodium chloride solution.

25 Instead of t-PA it is also possible to use the same amount of another plasminogen activator mentioned hereinbefore, such as scu-PA or a u-PA/t-PA hybrid plasminogen activator. Analogously, instead of desulfato hirudin HV1 it is also possible to use the same amount of another hirudin mentioned hereinbefore, such as hirudin PA or a mutant of desulfato hirudin HV1.

Claims

- 30 1. A pharmaceutical composition comprising in separate unit dosage forms as component A a plasminogen activator selected from the group consisting of u-PA and t-PA including glycosylated, partially glycosylated or unglycosylated forms thereof, a u-PA/t-PA hybrid plasminogen activator and mutants and fragments thereof retaining the fibrinolytic activity and as component B hirudin together with pharmaceutically acceptable carriers.
- 35 2. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator in the single chain form.
- 40 3. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator of the formula

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Ser Asn Glu Leu His Gln Val

Pro Ser Asn Cys Asp Cys Leu Asn Gly Gly Thr Cys Val Ser Asn
 5 Lys Tyr Phe Ser Asn Ile His Trp Cys Asn Cys Pro Lys Lys Phe
 Gly Gly Gln His Cys Glu Ile Asp Lys Ser Lys Thr Cys Tyr Glu
 Gly Asn Gly His Phe Tyr Arg Gly Lys Ala Ser Thr Asp Thr Met
 10 Gly Arg Pro Cys Leu Pro Trp Asn Ser Ala Thr Val Leu Gln Gln
 Thr Tyr His Ala His Arg Ser Asp Ala Leu Gln Leu Gly Leu Gly
 Lys His Asn Tyr Cys Arg Asn Pro Asp Asn Arg Arg Arg Pro Trp
 Cys Tyr Val Gln Val Gly Leu Lys Pro Leu Val Gln Glu Cys Met
 15 Val His Asp Cys Ala Asp Gly X₁ X₂ Pro Ser Ser Pro Pro Glu
 Glu Leu Lys Phe Gln Cys Gly Gln Lys Thr Leu Arg Pro Y₁ Y₂
 Y₃ Ile Ile Gly Gly Glu Phe Thr Thr Ile Glu Asn Gln Pro Trp
 20 Phe Ala Ala Ile Tyr Arg Arg His Arg Gly Gly Ser Val Thr Tyr
 Val Cys Gly Gly Ser Leu Ile Ser Pro Cys Trp Val Ile Ser Ala
 Thr His Cys Phe Ile Asp Tyr Pro Lys Lys Glu Asp Tyr Ile Val
 25 Tyr Leu Gly Arg Ser Arg Leu Asn Ser Asn Thr Gln Gly Glu Met
 Lys Phe Glu Val Glu Asn Leu Ile Leu His Lys Asp Tyr Ser Ala

Asp Thr Leu Ala His His Asn Asp Ile Ala Leu Leu Lys Ile Arg
 Ser Lys Glu Gly Arg Cys Ala Gln Pro Ser Arg Thr Ile Gln Thr
 Ile Cys Leu Pro Ser Met Tyr Asn Asp Pro Gln Phe Gly Thr Ser
 35 Cys Glu Ile Thr Gly Phe Gly Lys Glu Z₁ Ser Z₂ Asp Tyr Leu
 Tyr Pro Glu Gln Leu Lys Met Thr Val Val Lys Leu Ile Ser His
 Arg Glu Cys Gln Gln Pro His Tyr Tyr Gly Ser Glu Val Thr Thr
 40 Lys Met Leu Cys Ala Ala Asp Pro Gln Trp Lys Thr Asp Ser Cys
 Gln Gly Asp Ser Gly Gly Pro Leu Val Cys Ser Leu Gln Gly Arg
 Met Thr Leu Thr Gly Ile Val Ser Trp Gly Arg Gly Cys Ala Leu
 45 Lys Asp Lys Pro Gly Val Tyr Thr Arg Val Ser His Phe Leu Pro
 Trp Ile Arg Ser His Thr Lys Glu Glu Asn Gly Leu Ala Leu

(I)

in which X₁ and X₂ independently from each other represent Lys, an amino acid residue other than a
 basic amino acid residue or a chemical bond, Y₁ is Arg, an amino acid residue other than a basic
 amino acid residue or a chemical bond, Y₂ is Phe, an acidic amino acid residue or a chemical bond, Y₃
 55 is Lys, an amino acid residue other than a basic amino acid residue or a chemical bond, Z₁ is Asn or is
 another amino acid residue, and Z₂ is Thr or another amino acid residue different from Ser.



4. A pharmaceutical composition according to claim 3 comprising as component A a plasminogen activator of the formula I in which X_1 is Lys, Gly or Ser, X_2 represents Lys, Y_1 is Arg, Y_2 is Phe, Asp or Glu, Y_3 is Lys, Z_1 is Asn or Gln and Z_2 is Thr.
5. A pharmaceutical composition according to claim 3 comprising as component A a plasminogen activator selected from the group consisting of scu-PA, [Gly¹³⁵]-scu-PA, [Ser¹³⁵]-scu-PA, [Asp¹⁵⁷]-scu-PA, [Ser¹³⁵,Asp¹⁵⁷]-scu-PA and [Gly¹³⁵,Asp¹⁵⁷]-scu-PA wherein Asn³⁰² is yeast specifically glycosylated, and [Gln³⁰²]-scu-PA, [Gly¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA and [Ser¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA.
6. A pharmaceutical composition according to claim 3 comprising as component A scu-PA.
7. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator of the formula

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Ser Tyr Gln Val Ile Cys Arg Asp Glu Lys Thr Gln Met Ile Tyr Gln Gln
 His Gln Ser Trp Leu Arg Pro Val Leu Arg Ser Asn Arg Val Glu Tyr Cys
 5 Trp Cys Asn Ser Gly Arg Ala Gln Cys His Ser Val Pro Val Lys Ser Cys
 Ser Glu Pro Arg Cys Phe Asn Gly Gly Thr Cys Gln Gln Ala Leu Tyr Phe
 Ser Asp Phe Val Cys Gln Cys Pro Glu Gly Phe Ala Gly Lys Cys Cys Glu
 10 Ile Asp Thr Arg Ala Thr Cys Tyr Glu Asp Gln Gly Ile Ser Tyr Arg Gly
 Thr Trp Ser Thr Ala Glu Ser Gly Ala Glu Cys Thr Asn Trp X₁ Ser Y₁
 Ala Leu Ala Gln Lys Pro Tyr Ser Gly Arg Arg Pro Asp Ala Ile Arg Leu
 Gly Leu Gly Asn His Asn Tyr Cys Arg Asn Pro Asp Arg Asp Ser Lys Pro
 15 Trp Cys Tyr Val Phe Lys Ala Gly Lys Tyr Ser Ser Glu Phe Cys Ser Thr
 Pro Ala Cys Ser Glu Gly Asn Ser Asp Cys Tyr Phe Gly X₂ Gly Y₂ Ala
 Tyr Arg Gly Thr His Ser Leu Thr Glu Ser Gly Ala Ser Cys Leu Pro Trp
 20 Asn Ser Met Ile Leu Ile Gly Lys Val Tyr Thr Ala Gln Asn Pro Ser Ala
 Gln Ala Leu Gly Leu Gly Lys His Asn Tyr Cys Arg Asn Pro Asp Gly Asp
 Ala Lys Pro Trp Cys His Val Leu Lys Asn Arg Arg Leu Thr Trp Glu Tyr
 Cys Asp Val Pro Ser Cys Ser Thr Cys Gly Leu Arg Gln Tyr Ser Gln Pro
 25 Gln Phe Arg Ile Lys Gly Gly Leu Phe Ala Asp Ile Ala Ser His Pro Trp
 Gln Ala Ala Ile Phe Ala Lys His Arg Arg Ser Pro Gly Glu Arg Phe Leu
 Cys Gly Gly Ile Leu Ile Ser Ser Cys Trp Ile Leu Ser Ala Ala His Cys
 30 Phe Gln Glu Arg Phe Pro Pro His His Leu Thr Val Ile Leu Gly Arg Thr
 Tyr Arg Val Val Pro Gly Glu Glu Glu Gln Lys Phe Glu Val Glu Lys Tyr
 Ile Val His Lys Glu Phe Asp Asp Asp Thr Tyr Asp Asn Asp Ile Ala Leu
 35 Leu Gln Leu Lys Ser Asp Ser Ser Arg Cys Ala Gln Glu Ser Ser Val Val
 Arg Thr Val Cys Leu Pro Pro Ala Asp Leu Gln Leu Pro Asp Trp Thr Glu
 Cys Glu Leu Ser Gly Tyr Gly Lys His Glu Ala Leu Ser Pro Phe Tyr Ser
 40 Glu Arg Leu Lys Glu Ala His Val Arg Leu Tyr Pro Ser Ser Arg Cys Thr
 Ser Gln His Leu Leu X₃ Arg Y₃ Val Thr Asp Asn Met Leu Cys Ala Gly
 Asp Thr Arg Ser Gly Gly Pro Gln Ala Asn Leu His Asp Ala Cys Gln Gly
 Asp Ser Gly Gly Pro Leu Val Cys Leu Asn Asp Gly Arg Met Thr Leu Val
 45 Gly Ile Ile Ser Trp Gly Leu Gly Cys Gly Gln Lys Asp Val Pro Gly Val
 Tyr Thr Lys Val Thr Asn Tyr Leu Asp Trp Ile Arg Asp Asn Met Arg Pro

(II)

in which X₁, X₂ and X₃ each represent Asn or another genetically encoded amino acid, Y₁ and Y₂ each
 represent Ser or another genetically encoded amino acid other than Thr and Y₃ represents Thr or
 55 another genetically encoded amino acid other than Ser.

8. A pharmaceutical composition according to claim 7 comprising as component A a plasminogen activator of the formula II in which one, two or three of the radicals X₁, X₂ and X₃ represent Asn and



- the other(s) represent(s) an amino acid residue selected from Gln, Thr and Ser, Y₁ and Y₂ each represent Ser and Y₃ represents Thr, or in which X₁, X₂ and X₃ each represent Asn, one or two of the radicals Y₁ and Y₂ represent Ser and the other(s) is (are) selected from the group consisting of Ala and Asn and Y₃ represents Thr, or in which X₁, X₂ and X₃ each represent Asn, Y₁ and Y₂ each represent Ser and Y₃ is selected from the group consisting of Ala and Asn.
9. A pharmaceutical composition according to claim 7 comprising as component A a plasminogen activator selected from the group consisting of t-PA, [Asn¹¹⁹]-t-PA, [Ala¹⁸⁶]-t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹, Ala¹⁸⁶]-t-PA, [Asn¹¹⁹, Ala¹⁸⁶, Asn⁴⁵⁰]-t-PA and [Asn¹¹⁹, Ala¹⁸⁶, Ala⁴⁵⁰]-t-PA.
 10. A pharmaceutical composition according to claim 9 comprising as component A t-PA.
 11. A pharmaceutical composition according to claim 1 comprising as component A a single-chain hybrid plasminogen activator comprising an amino acid sequence containing all or discrete A-chain domains of human u-PA or discrete A-chain domains of human u-PA and human t-PA, linked in series to the catalytic region of human t-PA, and a single-chain hybrid PA comprising an amino acid sequence containing all or discrete A-chain domains of human t-PA or discrete A-chain domains of human t-PA and u-PA, linked in series to the catalytic region of human u-PA, or a single-chain hybrid plasminogen activator differing from u-PA in that the amino acid segment following the u-PA kringle up to the u-PA activation site is replaced by the amino acid segment following the t-PA kringle 2 up to the t-PA activation site.
 12. A pharmaceutical composition according to claim 11 comprising as component A a single-chain hybrid plasminogen activator comprising an amino acid sequence selected from the group consisting of an amino acid sequence containing all of the A-chain domains of human t-PA, an amino acid sequence containing discrete A-chain domains of human t-PA, such as the finger domain or a kringle, especially the kringle 2, domain of human t-PA, and an amino acid sequence containing two, three or four A-chain domains of human t-PA and/or human u-PA, especially two or three domains of human t-PA or two or three domains of human u-PA and human t-PA, such as the finger, growth factor and kringle 2 domains of human t-PA, the finger and kringle 2 domains of human t-PA or the u-PA growth factor and t-PA kringle 2 domains, which amino acid sequence is linked in series to the catalytic region of human u-PA, and an amino acid sequence containing the u-PA growth factor and t-PA kringle 2 domains which amino acid sequence is linked in series to the catalytic region of human t-PA.
 13. A pharmaceutical composition according to claim 11 comprising as component A a single-chain hybrid plasminogen activator selected from the group consisting of a hybrid plasminogen activator comprising the A-chain of u-PA or an A-chain essentially consisting of the u-PA growth factor and the t-PA kringle 2 domains linked in series to the catalytic region (B-chain) of t-PA, and a hybrid plasminogen activator comprising the A-chain of t-PA, an A-chain essentially consisting of the finger domain of t-PA, an A-chain essentially consisting of the u-PA growth factor and t-PA kringle 2 domains, an A-chain essentially consisting of the t-PA finger and kringle 2 domains or an A-chain essentially consisting of the t-PA finger, growth factor and kringle 2 domains said A-chain being linked in series to the catalytic region (B-chain) of u-PA, wherein the A-chain is linked to the B-chain via a junction sequence comprising an activation site and a cysteine residue capable of forming a sulphur-sulphur bond to the B-chain or the single chain hybrid plasminogen activator comprises an A-chain essentially consisting of the t-PA kringle 2 domain linked to the catalytic region (B-chain) of u-PA at the activation site.
 14. A pharmaceutical composition according to claim 11 comprising as component A a single chain hybrid plasminogen activator selected from the group consisting of such a hybrid PA comprising an A-chain essentially consisting of the u-PA growth factor domain and the t-PA kringle 2 domain linked in series to the catalytic region (B-chain) of t-PA, and a single-chain hybrid plasminogen activator comprising an A-chain essentially consisting of the t-PA kringle 2 domain or of the t-PA finger and kringle 2 domains linked in series to the catalytic region (B-chain) of u-PA, wherein the junction between the A-chain domain(s) and the B-chain is at the activation site.
 15. A pharmaceutical composition according to claim 11 comprising as component A a single-chain hybrid plasminogen activator selected from the group consisting of [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] and [tPA(1-3)-tPA(176-275)-uPA(159-411)].



16. A pharmaceutical composition according to claim 1 comprising as component A a mutant of a u-PA/t-PA hybrid plasminogen activator in which at least one of the N-glycosylation sites is modified such that glycosylation cannot take place at this site.

5 17. A pharmaceutical composition according to claim 1 comprising as component B a hirudin selected from the group consisting of hirudin variant HV1 having the formula

10 Val Val Tyr Thr Asp Cys Thr Glu Ser Gly
Gln Asn Leu Cys Leu Cys Glu Gly Ser Asn
Val Cys Gly Gln Gly Asn Lys Cys Ile Leu
Gly Ser Asp Gly Glu Lys Asn Gln Cys Val
15 Thr Gly Glu Gly Thr Pro Lys Pro Gln Ser
His Asn Asp Gly Asp Phe Glu Glu Ile Pro
Glu Glu ^W Tyr Leu Gln

(III)

wherein W represents the phenolic hydroxy group of Tyr (desulfato hirudin HV1) or a group of the formula -O-SO₃H (hirudin HV1), hirudin variant HV2 having the formula

25 Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
30 Ile Leu Gly Ser Asn Gly Lys Gly Asn Gln Cys Val Thr Gly
Glu Gly Thr Pro Asn Pro Glu Ser His Asn Asn Gly Asp Phe
Glu Glu Ile Pro Glu Glu Tyr Leu Gln

(IV)

and hirudin variant PA having the formula

40 Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
Ile Leu Gly Ser Gln Gly Lys Asp Asn Gln Cys Val Thr Gly
45 Glu Gly Thr Pro Lys Pro Gln Ser His Asn Gln Gly Asp Phe
Glu Pro Ile Pro Glu Asp Ala Tyr Asp Glu

(V)

or a mutant or a fragment of any of said hirudin variants retaining the antithrombotic activity.

55 18. A pharmaceutical composition according to claim 17 comprising as component B a hirudin selected from the group consisting of desulfato hirudin HV1, hirudin HV1, hirudin HV2 und hirudin PA.



19. A pharmaceutical composition according to claim 18 comprising as component B desulfato hirudin HV1.
20. A pharmaceutical composition according to claim 17 comprising as component B a hirudin selected from the group consisting of [Ile^{1,2}]-desulfato hirudin HV1, [Gly¹]-desulfato hirudin HV1, [Gln^{57,58,61,62}]-desulfato hirudin HV1, [Pro⁶⁶]-desulfato hirudin HV1, [Gln²⁷,Arg⁴⁷]-desulfato hirudin HV1, [Met⁰,Gln²⁷,Arg⁴⁷]-desulfato hirudin HV1, [Gln²⁷,Gln³⁶,Arg⁴⁷]-desulfato hirudin HV1, [des-Val¹,Thr²]-desulfato hirudin HV1 and [Lys⁴⁷]-hirudin HV2.
21. A pharmaceutical composition according to claim 1 comprising as component A a plasminogen activator selected from the group consisting of scu-PA, t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹, Ala¹⁸⁶, Ala⁴⁵⁰]-t-PA, [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] and [tPA(1-3)-tPA(176-275)-uPA(159-411)], and as component B desulfato hirudin HV1.
22. A pharmaceutical composition according to claim 1 for use in a method of treatment of the human or animal body.
23. A pharmaceutical composition according to claim 1 for use in the prevention or treatment of thrombosis or diseases caused by thrombosis, arteriosclerosis, myocardial and cerebral infarction, venous thrombosis, thromboembolism, post-surgical thrombosis and thrombophlebitis.
24. A pharmaceutical composition according to claim 1 for use in the prevention or treatment of myocardial infarction.
25. A pharmaceutical composition according to claim 1 wherein the ratio by weight of component A to component B is between 20:1 and 2:1.
26. A pharmaceutical composition according to claim 25 wherein the ratio by weight of component A to component B is between 5:1 and 2:1.

Patentansprüche

1. Pharmazeutische Zusammensetzung, die in getrennten Einheitsdosierungsformen als Komponente A einen Plasminogenaktivator ausgewählt aus der Gruppe bestehend aus u-PA und t-PA einschließlich der glycosylierten, partiell glycosylierten oder unglycosylierten Formen davon, einem u-PA/t-PA-Hybrid-Plasminogenaktivator und Mutanten und Fragmenten davon, die die fibrinolytische Aktivität beibehalten, und als Komponente B Hirudin zusammen mit pharmazeutisch unbedenklichen Trägern umfaßt.
2. Pharmazeutische Zusammensetzung nach Anspruch 1, die als Komponente A einen Plasminogenaktivator in der einkettigen Form enthält.
3. Pharmazeutische Zusammensetzung nach Anspruch 1, umfassend als Komponente A einen Plasminogenaktivator der Formel



Ser' Asn Glu Leu His Gln Val
 Pro Ser Asn Cys Asp Cys Leu Asn Gly Gly Thr Cys Val Ser Asn
 5 Lys Tyr Phe Ser Asn Ile His Trp Cys Asn Cys Pro Lys Lys Phe
 Gly Gly Gln His Cys Glu Ile Asp Lys Ser Lys Thr Cys Tyr Glu
 Gly Asn Gly His Phe Tyr Arg Gly Lys Ala Ser Thr Asp Thr Met
 10 Gly Arg Pro Cys Leu Pro Trp Asn Ser Ala Thr Val Leu Gln Gln
 Thr Tyr His Ala His Arg Ser Asp Ala Leu Gln Leu Gly Leu Gly
 Lys His Asn Tyr Cys Arg Asn Pro Asp Asn Arg Arg Arg Pro Trp
 Cys Tyr Val Gln Val Gly Leu Lys Pro Leu Val Gln Glu Cys Met
 15 Val His Asp Cys Ala Asp Gly X₁ X₂ Pro Ser Ser Pro Pro Glu
 Glu Leu Lys Phe Gln Cys Gly Gln Lys Thr Leu Arg Pro Y₁ Y₂
 Y₃ Ile Ile Gly Gly Glu Phe Thr Thr Ile Glu Asn Gln Pro Trp
 20 Phe Ala Ala Ile Tyr Arg Arg His Arg Gly Gly Ser Val Thr Tyr
 Val Cys Gly Gly Ser Leu Ile Ser Pro Cys Trp Val Ile Ser Ala
 Thr His Cys Phe Ile Asp Tyr Pro Lys Lys Glu Asp Tyr Ile Val
 25 Tyr Leu Gly Arg Ser Arg Leu Asn Ser Asn Thr Gln Gly Glu Met
 Lys Phe Glu Val Glu Asn Leu Ile Leu His Lys Asp Tyr Ser Ala
 Asp Thr Leu Ala His His Asn Asp Ile Ala Leu Leu Lys Ile Arg
 30 Ser Lys Glu Gly Arg Cys Ala Gln Pro Ser Arg Thr Ile Gln Thr
 Ile Cys Leu Pro Ser Met Tyr Asn Asp Pro Gln Phe Gly Thr Ser
 Cys Glu Ile Thr Gly Phe Gly Lys Glu Z₁ Ser Z₂ Asp Tyr Leu
 35 Tyr Pro Glu Gln Leu Lys Met Thr Val Val Lys Leu Ile Ser His
 Arg Glu Cys Gln Gln Pro His Tyr Tyr Gly Ser Glu Val Thr Thr
 40 Lys Met Leu Cys Ala Ala Asp Pro Gln Trp Lys Thr Asp Ser Cys
 Gln Gly Asp Ser Gly Gly Pro Leu Val Cys Ser Leu Gln Gly Arg
 Met Thr Leu Thr Gly Ile Val Ser Trp Gly Arg Gly Cys Ala Leu
 45 Lys Asp Lys Pro Gly Val Tyr Thr Arg Val Ser His Phe Leu Pro
 Trp Ile Arg Ser His Thr Lys Glu Glu Asn Gly Leu Ala Leu

(I)

worin X₁ und X₂ unabhängig voneinander Lys, einen anderen als einen basischen Aminosäurerest oder
 eine chemische Bindung darstellen, Y₁ Arg, ein anderer als ein basischer Aminosäurerest oder eine
 chemische Bindung ist, Y₂ Phe, ein saurer Aminosäurerest oder eine chemische Bindung ist, Y₃ Lys,
 55 ein anderer als ein basischer Aminosäurerest oder eine chemische Bindung ist, Z₁ Asn oder ein
 anderer Aminosäurerest ist und Z₂ Thr oder ein anderer, von Ser verschiedener Aminosäurerest ist.



4. Pharmazeutische Zusammensetzung nach Anspruch 3, die als Komponente A einen Plasminogenaktivator der Formel I enthält, worin X₁ Lys, Gly oder Ser ist, X₂ Lys, Y₁ Arg, Y₂ Phe, Asp oder Glu ist, Y₃ Lys ist, Z₁ Asn oder Gln ist und Z₂ Thr ist.
5. Pharmazeutische Zusammensetzung nach Anspruch 3, die als Komponente A einen Plasminogenaktivator ausgewählt aus der Gruppe bestehend aus scu-PA, [Gly¹³⁵]-scu-PA, [Ser¹³⁵]-scu-PA, [Asp¹⁵⁷]-scu-PA, [Ser¹³⁵,Asp¹⁵⁷]-scu-PA und [Gly¹³⁵,Asp¹⁵⁷]-scu-PA, worin Asn³⁰² Hefespezifisch glycosyliert ist, und [Gln³⁰²]-scu-PA, [Gly¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA und [Ser¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA enthält.
6. Pharmazeutische Zusammensetzung nach Anspruch 3, die als Komponente A scu-PA enthält.
7. Pharmazeutische Zusammensetzung nach Anspruch 1, die als Komponente A einen Plasminogenaktivator der Formel

15

Ser Tyr Gln Val Ile Cys Arg Asp Glu Lys Thr Gln Met Ile Tyr Gln Gln
 His Gln Ser Trp Leu Arg Pro Val Leu Arg Ser Asn Arg Val Glu Tyr Cys
 Trp Cys Asn Ser Gly Arg Ala Gln Cys His Ser Val Pro Val Lys Ser Cys
 Ser Glu Pro Arg Cys Phe Asn Gly Gly Thr Cys Gln Gln Ala Leu Tyr Phe
 Ser Asp Phe Val Cys Gln Cys Pro Glu Gly Phe Ala Gly Lys Cys Cys Glu
 Ile Asp Thr Arg Ala Thr Cys Tyr Glu Asp Gln Gly Ile Ser Tyr Arg Gly
 Thr Trp Ser Thr Ala Glu Ser Gly Ala Glu Cys Thr Asn Trp X₁ Ser Y₁
 Ala Leu Ala Gln Lys Pro Tyr Ser Gly Arg Arg Pro Asp Ala Ile Arg Leu
 Gly Leu Gly Asn His Asn Tyr Cys Arg Asn Pro Asp Arg Asp Ser Lys Pro
 Trp Cys Tyr Val Phe Lys Ala Gly Lys Tyr Ser Ser Glu Phe Cys Ser Thr
 Pro Ala Cys Ser Glu Gly Asn Ser Asp Cys Tyr Phe Gly X₂ Gly Y₂ Ala
 Tyr Arg Gly Thr His Ser Leu Thr Glu Ser Gly Ala Ser Cys Leu Pro Trp
 Asn Ser Met Ile Leu Ile Gly Lys Val Tyr Thr Ala Gln Asn Pro Ser Ala

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Gln Ala Leu Gly Leu Gly Lys His Asn Tyr Cys Arg Asn Pro Asp Gly Asp
 Ala Lys Pro Trp Cys His Val Leu Lys Asn Arg Arg Leu Thr Trp Glu Tyr
 Cys Asp Val Pro Ser Cys Ser Thr Cys Gly Leu Arg Gln Tyr Ser Gln Pro
 5 Gln Phe Arg Ile Lys Gly Gly Leu Phe Ala Asp Ile Ala Ser His Pro Trp
 Gln Ala Ala Ile Phe Ala Lys His Arg Arg Ser Pro Gly Glu Arg Phe Leu
 Cys Gly Gly Ile Leu Ile Ser Ser Cys Trp Ile Leu Ser Ala Ala His Cys
 10 Phe Gln Glu Arg Phe Pro Pro His His Leu Thr Val Ile Leu Gly Arg Thr
 Tyr Arg Val Val Pro Gly Glu Glu Glu Gln Lys Phe Glu Val Glu Lys Tyr
 Ile Val His Lys Glu Phe Asp Asp Asp Thr Tyr Asp Asn Asp Ile Ala Leu
 15 Leu Gln Leu Lys Ser Asp Ser Ser Arg Cys Ala Gln Glu Ser Ser Val Val
 Arg Thr Val Cys Leu Pro Pro Ala Asp Leu Gln Leu Pro Asp Trp Thr Glu
 Cys Glu Leu Ser Gly Tyr Gly Lys His Glu Ala Leu Ser Pro Phe Tyr Ser
 20 Glu Arg Leu Lys Glu Ala His Val Arg Leu Tyr Pro Ser Ser Arg Cys Thr
 Ser Gln His Leu Leu X₁ Arg Y₁ Val Thr Asp Asn Met Leu Cys Ala Gly
 Asp Thr Arg Ser Gly Gly Pro Gln Ala Asn Leu His Asp Ala Cys Gln Gly
 Asp Ser Gly Gly Pro Leu Val Cys Leu Asn Asp Gly Arg Met Thr Leu Val
 25 Gly Ile Ile Ser Trp Gly Leu Gly Cys Gly Gln Lys Asp Val Pro Gly Val
 Tyr Thr Lys Val Thr Asn Tyr Leu Asp Trp Ile Arg Asp Asn Met Arg Pro

(II)

enthält, worin X₁, X₂ und X₃ jedes Asn oder eine andere genetisch codierte Aminosäure darstellen, Y₁ und Y₂ jedes Ser oder eine andere genetisch codierte Aminosäure außer Thr darstellen und Y₃ Thr oder eine andere genetisch codierte Aminosäure außer Ser darstellt.

8. Pharmazeutische Zusammensetzung nach Anspruch 7, umfassend als Komponente A einen Plasminogenaktivator der Formel II, worin einer, zwei oder drei der Reste X₁, X₂ und X₃ Asn darstellen und der(die) andere(n) einen aus Gln, Thr und Ser ausgewählten Aminosäurerest darstellt(darstellen), Y₁ und Y₂ jedes Ser darstellt und Y₃ Thr darstellt, oder worin X₁, X₂ und X₃ jedes Asn darstellen, einer oder zwei der Reste Y₁ und Y₂ Ser darstellen und der(die) andere(n) aus der aus Ala und Asn bestehenden Gruppe ausgewählt ist(sind) und Y₃ Thr darstellt, oder worin X₁, X₂ und X₃ jedes Asn darstellen, Y₁ und Y₂ jedes Ser darstellt und Y₃ aus der aus Ala und Asn bestehenden Gruppe ausgewählt ist.
9. Pharmazeutische Zusammensetzung nach Anspruch 7, die als Komponente A einen Plasminogenaktivator, der ausgewählt ist aus der Gruppe bestehend aus t-PA, [Asn¹¹⁹]-t-PA, [Ala¹⁸⁶]-t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹,Ala¹⁸⁶]-t-PA, [Asn¹¹⁹,Ala¹⁸⁶,Asn⁴⁵⁰]-t-PA und [Asn¹¹⁹,Ala¹⁸⁶,Ala⁴⁵⁰]-t-PA, enthält.
10. Pharmazeutische Zusammensetzung nach Anspruch 9, die als Komponente A t-PA enthält.
11. Pharmazeutische Zusammensetzung nach Anspruch 1, die als Komponente A einen einkettigen Hybrid-Plasminogenaktivator enthält, umfassend eine Aminosäuresequenz, die alle oder diskrete A-Ketten-Domänen von Human-u-PA oder diskrete A-Ketten-Domänen von Human-u-PA und Human-t-PA enthält, der Reihe nach verbunden mit der katalytischen Region von Human-t-PA, und einen einkettigen Hybrid-PA, umfassend eine Aminosäuresequenz, die alle oder diskrete A-Ketten-Domänen von Human-t-PA oder diskrete A-Ketten-Domänen von Human-t-PA und u-PA enthält, der Reihe nach verbunden mit der katalytischen Region von Human-u-PA, oder einen einkettigen Hybrid-Plasminogenaktivator, der sich von u-PA dadurch unterscheidet, daß das dem u-PA-Kringel folgende Aminosäuresegment bis zur u-PA-Aktivierungsstelle durch das dem t-PA-Kringel 2 folgende Aminosäuresegment bis zur t-PA-



Aktivierungsstelle ersetzt ist.

- 5 12. Pharmazeutische Zusammensetzung nach Anspruch 11, umfassend als Komponente A einen einkettigen Hybrid-Plasminogenaktivator, umfassend eine Aminosäuresequenz, die ausgewählt ist aus der Gruppe bestehend aus einer Aminosäuresequenz, die alle A-Ketten-Domänen von Human-t-PA enthält, einer Aminosäuresequenz, die diskrete A-Ketten-Domänen von Human-t-PA enthält, wie die Fingerdomäne oder eine Kringel-, insbesondere Kringel 2-Domäne von Human-t-PA, und einer Aminosäuresequenz, die zwei, drei oder vier A-Ketten-Domänen von Human-t-PA und/oder Human-u-PA enthält, insbesondere zwei oder drei Domänen von Human-t-PA oder zwei oder drei Domänen von Human-u-PA und Human-t-PA, wie die Finger-, Wachstumsfaktor- und Kringel 2-Domänen von Human-t-PA, die Finger- und Kringel 2-Domänen von Human-t-PA oder die u-PA-Wachstumsfaktor- und t-PA-Kringel 2-Domänen, welche Aminosäuresequenz der Reihe nach verbunden ist mit der katalytischen Region von Human-u-PA, und einer Aminosäuresequenz, die die u-PA-Wachstumsfaktor- und t-PA-Kringel 2-Domänen enthält, welche Aminosäuresequenz der Reihe nach mit der katalytischen Region von Human-t-PA verbunden ist.
- 10 13. Pharmazeutische Zusammensetzung nach Anspruch 11, umfassend als Komponente A einen einkettigen Hybrid-Plasminogenaktivator, ausgewählt aus der Gruppe bestehend aus einem Hybrid-Plasminogenaktivator, umfassend die A-Kette von u-PA oder eine A-Kette, bestehend im wesentlichen aus den u-PA-Wachstumsfaktor- und den t-PA-Kringel 2-Domänen, der Reihe nach verbunden mit der katalytischen Region (B-Kette) von t-PA, und einem Hybrid-Plasminogenaktivator, umfassend die A-Kette von t-PA, eine A-Kette, die im wesentlichen aus der Fingerdomäne von t-PA besteht, eine A-Kette, die im wesentlichen aus den u-PA-Wachstumsfaktor- und t-PA-Kringel 2-Domänen besteht, eine A-Kette, die im wesentlichen aus den t-PA-Finger- und -Kringel 2-Domänen besteht, oder eine A-Kette, die im wesentlichen aus den t-PA-Finger-, -Wachstumsfaktor- und -Kringel 2-Domänen besteht, wobei die A-Kette der Reihe nach mit der katalytischen Region (B-Kette) von u-PA verbunden ist, worin die A-Kette mit der B-Kette über eine Verbindungssequenz verbunden ist, die eine Aktivierungsstelle und einen Cysteinrest enthält, der befähigt ist, mit der B-Kette eine Schwefel-Schwefel-Bindung zu bilden, oder der einkettige Hybrid-Plasminogenaktivator umfaßt eine A-Kette, die im wesentlichen aus der t-PA-Kringel 2-Domäne besteht, die verbunden ist mit der katalytischen Region (B-Kette) von u-PA an der Aktivierungsstelle.
- 20 14. Pharmazeutische Zusammensetzung nach Anspruch 11, umfassend als Komponente A einen einkettigen Hybrid-Plasminogenaktivator, ausgewählt aus der Gruppe bestehend aus einem Hybrid-PA umfassend eine A-Kette, bestehend im wesentlichen aus der u-PA-Wachstumsfaktor-Domäne und der t-PA-Kringel 2-Domäne, der Reihe nach verbunden mit der katalytischen Region (B-Kette) von t-PA, und einem einkettigen Hybrid-Plasminogenaktivator, umfassend eine A-Kette, bestehend im wesentlichen aus der t-PA-Kringel 2-Domäne oder den t-PA-Finger- und -Kringel 2-Domänen, der Reihe nach verbunden mit der katalytischen Region (B-Kette) von u-PA, worin die Verbindung zwischen der(den) A-Ketten-Domäne(n) und der B-Kette an der Aktivierungsstelle ist.
- 25 15. Pharmazeutische Zusammensetzung nach Anspruch 11, umfassend als Komponente A einen einkettigen Hybrid-Plasminogenaktivator, ausgewählt aus der Gruppe bestehend aus [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] und [tPA(1-3)-tPA(176-275)-uPA(159-411)].
- 30 16. Pharmazeutische Zusammensetzung nach Anspruch 1, umfassend als Komponente A eine Mutante eines u-PA/t-PA-Hybrid-Plasminogenaktivators, worin zumindest einer der N-Glycosylierungsorte so modifiziert ist, daß Glycosylierung an diesem Ort nicht stattfinden kann.
- 35 17. Pharmazeutische Zusammensetzung nach Anspruch 1, die als Komponente B ein Hirudin enthält, ausgewählt aus der Gruppe bestehend aus Hirudin-Variante HV1 mit der Formel
- 40
- 45
- 50
- 55



5 Val Val Tyr Thr Asp Cys Thr Glu Ser Gly
 Gln Asn Leu Cys Leu Cys Glu Gly Ser Asn
 Val Cys Gly Gln Gly Asn Lys Cys Ile Leu
 Gly Ser Asp Gly Glu Lys Asn Gln Cys Val
 Thr Gly Glu Gly Thr Pro Lys Pro Gln Ser
 10 His Asn Asp Gly Asp Phe Glu Glu Ile Pro
 Glu Glu ^W Tyr Leu Gln

(III)

15 worin W die phenolische Hydroxygruppe von Tyr (Desulfatohirudin HV1) oder eine Gruppe der Formel
 -O-SO₃H (Hirudin HV1) darstellt, Hirudin-Variante HV2 mit der Formel

20 Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 Ile Leu Gly Ser Asn Gly Lys Gly Asn Gln Cys Val Thr Gly
 Glu Gly Thr Pro Asn Pro Glu Ser His Asn Asn Gly Asp Phe
 25 Glu Glu Ile Pro Glu Glu Tyr Leu Gln

(IV)

30 und Hirudin-Variante PA mit der Formel

35 Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 Ile Leu Gly Ser Gln Gly Lys Asp Asn Gln Cys Val Thr Gly
 Glu Gly Thr Pro Lys Pro Gln Ser His Asn Gln Gly Asp Phe
 40 Glu Pro Ile Pro Glu Asp Ala Tyr Asp Glu

(V)

45 oder einer Mutante oder einem Fragment einer dieser Hirudinvarianten, die die antithrombolische
 Wirkung beibehalten.

18. Pharmazeutische Zusammensetzung nach Anspruch 17, die als Komponente B ein Hirudin enthält,
 ausgewählt aus der Gruppe bestehend aus Desulfatohirudin HV1, Hirudin HV1, Hirudin HV2 und Hirudin
 50 PA.

19. Pharmazeutische Zusammensetzung nach Anspruch 18, die als Komponente B Desulfatohirudin HV1
 enthält.

55 20. Pharmazeutische Zusammensetzung nach Anspruch 17, die als Komponente B ein Hirudin enthält,
 ausgewählt aus der Gruppe bestehend aus [Ile^{1,2}]-Desulfatohirudin HV1, [Gly¹]-Desulfatohirudin HV1,
 [Gln^{57,58,61,62}]-Desulfatohirudin HV1, [Pro⁵⁶]-Desulfatohirudin HV1, [Gln²⁷, Arg⁴⁷]-Desulfatohirudin HV1,
 (Met⁰, Gln²⁷, Arg⁴⁷)-Desulfatohirudin HV1, [Gln²⁷, Gln³⁶, Arg⁴⁷]-Desulfatohirudin HV1, [des-Val¹, Thr²]-De-



sulfatohirudin HV1 und [Lys⁴⁷]-Hirudin HV2.

21. Pharmazeutische Zusammensetzung nach Anspruch 1, die als Komponente A einen Plasminogenaktivator, ausgewählt aus der Gruppe bestehend aus scu-PA, t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁵,Ala¹⁸⁶,Ala⁴⁵⁰]-t-PA, [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] und [t-PA(1-3)-tPA(176-275)-uPA(159-411)], und als Komponente B Desulfatohirudin HV1 enthält.
22. Pharmazeutische Zusammensetzung nach Anspruch 1 zur Verwendung bei einer Methode zur Behandlung des menschlichen oder tierischen Körpers.
23. Pharmazeutische Zusammensetzung nach Anspruch 1 zur Verwendung bei der Verhinderung oder Behandlung von Thrombose oder durch Thrombose hervorgerufenen Erkrankungen, Arteriosklerose, Myocardinfarkt, Hirninfarkt, venöser Thrombose, Thromboembolie, Thrombose nach chirurgischem Eingriff und Thrombophlebitis.
24. Pharmazeutische Zusammensetzung nach Anspruch 1 zur Verwendung beim Verhindern oder Behandeln von Myocardinfarkt.
25. Pharmazeutische Zusammensetzung nach Anspruch 1, worin das Gewichtsverhältnis der Komponente A zur Komponente B zwischen 20:1 und 2:1 liegt.
26. Pharmazeutische Zusammensetzung nach Anspruch 25, worin das Gewichtsverhältnis der Komponente A zur Komponente B zwischen 5:1 und 2:1 liegt.

25 Revendications

1. Composition pharmaceutique comprenant sous des formes de dosage unitaire séparées en tant que composant A un activateur du plasminogène
choisi parmi l'u-PA et le t-PA, y compris leurs formes glycosylées, partiellement glycosylées ou non-glycosylées, un activateur du plasminogène hybride u-PA/t-PA et leurs mutants et variantes conservant l'activité fibrinolytique et en tant que composant B de l'hirudine, conjointement avec des supports pharmaceutiquement acceptables.
2. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un activateur du plasminogène sous une forme à simple chaîne.
3. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un activateur du plasminogène de formule



Ser Asn Glu Leu His Gln Val
 Pro Ser Asn Cys Asp Cys Leu Asn Gly Gly Thr Cys Val Ser Asn
 Lys Tyr Phe Ser Asn Ile His Trp Cys Asn Cys Pro Lys Lys Phe
 Gly Gly Gln His Cys Glu Ile Asp Lys Ser Lys Thr Cys Tyr Glu
 Gly Asn Gly His Phe Tyr Arg Gly Lys Ala Ser Thr Asp Thr Met
 Gly Arg Pro Cys Leu Pro Trp Asn Ser Ala Thr Val Leu Gln Gln
 Thr Tyr His Ala His Arg Ser Asp Ala Leu Gln Leu Gly Leu Gly
 Lys His Asn Tyr Cys Arg Asn Pro Asp Asn Arg Arg Arg Pro Trp
 Cys Tyr Val Gln Val Gly Leu Lys Pro Leu Val Gln Glu Cys Met
 Val His Asp Cys Ala Asp Gly X₁ X₂ Pro Ser Ser Pro Pro Glu
 Glu Leu Lys Phe Gln Cys Gly Gln Lys Thr Leu Arg Pro Y₁ Y₂
 Y₃ Ile Ile Gly Gly Glu Phe Thr Thr Ile Glu Asn Gln Pro Trp
 Phe Ala Ala Ile Tyr Arg Arg His Arg Gly Gly Ser Val Thr Tyr
 Val Cys Gly Gly Ser Leu Ile Ser Pro Cys Trp Val Ile Ser Ala

Thr His Cys Phe Ile Asp Tyr Pro Lys Lys Glu Asp Tyr Ile Val
 Tyr Leu Gly Arg Ser Arg Leu Asn Ser Asn Thr Gln Gly Glu Met
 Lys Phe Glu Val Glu Asn Leu Ile Leu His Lys Asp Tyr Ser Ala

Asp Thr Leu Ala His His Asn Asp Ile Ala Leu Leu Lys Ile Arg
 Ser Lys Glu Gly Arg Cys Ala Gln Pro Ser Arg Thr Ile Gln Thr
 Ile Cys Leu Pro Ser Met Tyr Asn Asp Pro Gln Phe Gly Thr Ser
 Cys Glu Ile Thr Gly Phe Gly Lys Glu Z₁ Ser Z₂ Asp Tyr Leu
 Tyr Pro Glu Gln Leu Lys Met Thr Val Val Lys Leu Ile Ser His
 Arg Glu Cys Gln Gln Pro His Tyr Tyr Gly Ser Glu Val Thr Thr
 Lys Met Leu Cys Ala Ala Asp Pro Gln Trp Lys Thr Asp Ser Cys
 Gln Gly Asp Ser Gly Gly Pro Leu Val Cys Ser Leu Gln Gly Arg
 Met Thr Leu Thr Gly Ile Val Ser Trp Gly Arg Gly Cys Ala Leu
 Lys Asp Lys Pro Gly Val Tyr Thr Arg Val Ser His Phe Leu Pro
 Trp Ile Arg Ser His Thr Lys Glu Glu Asn Gly Leu Ala Leu

(I)

où X₁ et X₂ représentent, indépendamment l'un de l'autre, Lys, un résidu d'acide aminé autre qu'un
 résidu d'acide aminé basique ou une liaison chimique, Y₁ est Arg, un résidu d'acide aminé autre qu'un
 résidu d'acide aminé basique ou une liaison chimique, Y₂ est Phe, un résidu d'acide aminé acide ou
 une liaison chimique, Y₃ est Lys, un résidu d'acide aminé autre qu'un résidu d'acide aminé basique ou
 une liaison chimique, Z₁ est Asn ou est un autre résidu d'acide aminé, et Z₂ est Thr ou un autre résidu
 d'acide aminé différent de Ser.



4. Composition pharmaceutique selon la revendication 3 comprenant en tant que composant A un activateur du plasminogène de formule I dans laquelle X₁ est Lys, Gly ou Ser, X₂ représente Lys, X₁ est Arg, Y₂ est Phe, Asp ou Glu, Y₃ est Lys, Z₁ est Asn ou Gln et Z₂ est Thr.
5. Composition pharmaceutique selon la revendication 3 comprenant en tant que composant A un activateur du plasminogène choisi parmi scu-PA, [Gly¹³⁵]-scu-PA, [Ser¹³⁵]-scu-PA, [Asp¹⁵⁷]-scu-PA, [Ser¹³⁵,Asp¹⁵⁷]-scu-PA et [Gly¹³⁵,Asp¹⁵⁷]-scu-PA, où Asn³⁰² est glycosylé spécifiquement par une levure, et [Gln³⁰²]-scu-PA, [Gly¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA et [Ser¹³⁵,Asp¹⁵⁷,Gln³⁰²]-scu-PA.
6. Composition pharmaceutique selon la revendication 3, comprenant en tant que composant A scu-PA.
7. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un activateur du plasminogène de formule

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Ser Tyr Gln Val Ile Cys Arg Asp Glu Lys Thr Gln Met Ile Tyr Gln Gln
 His Gln Ser Trp Leu Arg Pro Val Leu Arg Ser Asn Arg Val Glu Tyr Cys
 Trp Cys Asn Ser Gly Arg Ala Gln Cys His Ser Val Pro Val Lys Ser Cys
 Ser Glu Pro Arg Cys Phe Asn Gly Gly Thr Cys Gln Gln Ala Leu Tyr Phe
 Ser Asp Phe Val Cys Gln Cys Pro Glu Gly Phe Ala Gly Lys Cys Cys Glu
 Ile Asp Thr Arg Ala Thr Cys Tyr Glu Asp Gln Gly Ile Ser Tyr Arg Gly
 Thr Trp Ser Thr Ala Glu Ser Gly Ala Glu Cys Thr Asn Trp X₁ Ser Y₁

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Ala Leu Ala Gln Lys Pro Tyr Ser Gly Arg Arg Pro Asp Ala Ile Arg Leu
 Gly Leu Gly Asn His Asn Tyr Cys Arg Asn Pro Asp Arg Asp Ser Lys Pro
 Trp Cys Tyr Val Phe Lys Ala Gly Lys Tyr Ser Ser Glu Phe Cys Ser Thr
 Pro Ala Cys Ser Glu Gly Asn Ser Asp Cys Tyr Phe Gly X₂ Gly Y₂ Ala
 Tyr Arg Gly Thr His Ser Leu Thr Glu Ser Gly Ala Ser Cys Leu Pro Trp
 Asn Ser Met Ile Leu Ile Gly Lys Val Tyr Thr Ala Gln Asn Pro Ser Ala

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Gln Ala Leu Gly Leu Gly Lys His Asn Tyr Cys Arg Asn Pro Asp Gly Asp
 Ala Lys Pro Trp Cys His Val Leu Lys Asn Arg Arg Leu Thr Trp Glu Tyr
 Cys Asp Val Pro Ser Cys Ser Thr Cys Gly Leu Arg Gln Tyr Ser Gln Pro

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Gln Phe Arg Ile Lys Gly Gly Leu Phe Ala Asp Ile Ala Ser His Pro Trp
 Gln Ala Ala Ile Phe Ala Lys His Arg Arg Ser Pro Gly Glu Arg Phe Leu
 Cys Gly Gly Ile Leu Ile Ser Ser Cys Trp Ile Leu Ser Ala Ala His Cys

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Phe Gln Glu Arg Phe Pro Pro His His Leu Thr Val Ile Leu Gly Arg Thr
 Tyr Arg Val Val Pro Gly Glu Glu Glu Gln Lys Phe Glu Val Glu Lys Tyr
 Ile Val His Lys Glu Phe Asp Asp Asp Thr Tyr Asp Asn Asp Ile Ala Leu
 Leu Gln Leu Lys Ser Asp Ser Ser Arg Cys Ala Gln Glu Ser Ser Val Val

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Arg Thr Val Cys Leu Pro Pro Ala Asp Leu Gln Leu Pro Asp Trp Thr Glu
 Cys Glu Leu Ser Gly Tyr Gly Lys His Glu Ala Leu Ser Pro Phe Tyr Ser
 5 Glu Arg Leu Lys Glu Ala His Val Arg Leu Tyr Pro Ser Ser Arg Cys Thr
 Ser Gln His Leu Leu X₃ Arg Y₃ Val Thr Asp Asn Met Leu Cys Ala Gly
 Asp Thr Arg Ser Gly Gly Pro Gln Ala Asn Leu His Asp Ala Cys Gln Gly
 10 Asp Ser Gly Gly Pro Leu Val Cys Leu Asn Asp Gly Arg Met Thr Leu Val
 Gly Ile Ile Ser Trp Gly Leu Gly Cys Gly Gln Lys Asp Val Pro Gly Val
 Tyr Thr Lys Val Thr Asn Tyr Leu Asp Trp Ile Arg Asp Asn Met Arg Pro

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(II)

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où X₁, X₂ et X₃ représentent chacun Asn ou un autre acide aminé codé génétiquement, Y₁ et Y₂ représentent chacun Ser ou un autre acide aminé codé génétiquement autre que Thr et Y₃ représente Thr ou un autre acide aminé codé génétiquement autre que Ser.

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8. Composition pharmaceutique selon la revendication 7, comprenant en tant que composant A un activateur du plasminogène de formule II dans laquelle un, deux ou trois des radicaux X₁, X₂ et X₃ représente(nt) Asn et le ou les autres représente(nt) un résidu d'acide aminé choisi parmi Gln, Thr et Ser, Y₁ et X₂ représentent chacun Ser et Y₃ représente Thr, ou dans laquelle X₁, X₂ et X₃ représentent chacun Asn, un ou deux des radicaux Y₁ et Y₂ représente(nt) Ser et le ou les autres est(sont) choisi(s) dans le groupe composé de Ala et Asn et Y₃ représente Thr, ou dans laquelle X₁, X₂ et X₃ représentent chacun Asn, Y₁ et Y₂ représentent chacun Ser et Y₃ est choisi parmi Ala et Asn.

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9. Composition pharmaceutique selon la revendication 7, comprenant en tant que composant A un activateur du plasminogène choisi parmi t-PA, [Asn¹¹⁹]-t-PA, [Ala¹⁸⁶]-t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹,Ala¹⁸⁶]-t-PA, [Asn¹¹⁹,Ala¹⁸⁶,Asn⁴⁵⁰]-t-PA et [Asn¹¹⁹,Ala¹⁸⁶,Ala⁴⁵⁰]-t-PA

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10. Composition pharmaceutique selon la revendication 9, comprenant en tant que composant A t-PA.

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11. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un activateur du plasminogène hybride à simple chaîne comprenant une séquence d'acides aminés contenant la totalité ou certains des domaines de chaîne A d'u-PA humain ou certains domaines de chaîne A d'u-PA humain et de t-PA humain, liés en série à la région catalytique du t-PA humain, et un PA hybride à simple chaîne comprenant une séquence d'acides aminés contenant la totalité ou certains des domaines de chaîne A du t-PA humain ou certains domaines de chaîne A de t-PA humain et d'u-PA humain, liés en série à la région catalytique de l'u-PA humain, ou un activateur du plasminogène hybride à simple chaîne différent de l'u-PA en ce sens que le segment d'acide aminé suivant la boucle de l'u-PA jusqu'au site d'activation de l'u-PA est remplacé par le segment d'acide aminé suivant la boucle 2 du t-PA jusqu'au site d'activation du t-PA.

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12. Composition pharmaceutique selon la revendication 11, comprenant en tant que composant A un activateur du plasminogène hybride à simple chaîne comprenant une séquence d'acides aminés choisie dans le groupe consistant en une séquence d'acides aminés contenant tous les domaines de chaîne A du t-PA humain, une séquence d'acides aminés contenant des domaines distincts de chaîne A de t-PA humain, tels que le domaine du doigt ou un domaine de la boucle, en particulier la boucle 2, de t-PA humain, et une séquence d'acides aminés contenant deux, trois ou quatre domaines de chaîne A de t-PA humain et/ou d'u-PA humain, en particulier deux ou trois domaines de t-PA humain ou deux ou trois domaines d'u-PA humain et de t-PA humain, tels que les domaines du doigt, du facteur de croissance et de la boucle 2 du t-PA humain, les domaines du doigt et de la boucle 2 du t-PA humain ou les domaines du facteur de croissance de l'u-PA et de la boucle 2 du t-PA, laquelle séquence d'acides aminés est liée en série à la région catalytique de l'u-PA humain, et une séquence d'acides aminés contenant les domaines du facteur de croissance d'u-PA et de la boucle 2 de t-PA, laquelle



séquence d'acides aminés est liée en série à la région catalytique du t-PA humain.

- 5 13. Composition pharmaceutique selon la revendication 11, comprenant en tant que composant A un activateur du plasminogène hybride à simple chaîne choisi dans le groupe consistant en un activateur du plasminogène hybride comprenant la chaîne A de l'u-PA ou une chaîne A consistant essentiellement en les domaines du facteur de croissance de l'u-PA et de la boucle 2 du t-PA, liés en série à la région catalytique (chaîne B) du t-PA, et un activateur du plasminogène hybride comprenant la chaîne A du t-PA, une chaîne A consistant essentiellement en le domaine du doigt du t-PA, une chaîne A consistant essentiellement en les domaines du facteur de croissance de l'u-PA et de la boucle 2 du t-PA, une chaîne A consistant essentiellement en les domaines du doigt et de la boucle 2 du t-PA ou une chaîne A consistant essentiellement en les domaines du doigt, du facteur de croissance et de la boucle 2 du t-PA, la dite chaîne étant liée en série à la région catalytique (chaîne B) de l'u-PA, où la chaîne A est liée à la chaîne B via une séquence de jonction comprenant un site d'activation et un résidu de cystéine apte à former une liaison soufre-soufre avec la chaîne B ou l'activateur du plasminogène hybride à simple chaîne comprend une chaîne A consistant essentiellement en le domaine de la boucle 2 du t-PA relié à la région catalytique (chaîne B) de l'u-PA sur le site d'activation.
- 10 14. Composition pharmaceutique selon la revendication 11, comprenant en tant que composant A un activateur du plasminogène hybride à simple chaîne choisi dans le groupe consistant en un tel PA hybride comprenant une chaîne A consistant essentiellement en le domaine du facteur de croissance de l'u-PA et le domaine de la boucle 2 du t-PA liés en série à la région catalytique (chaîne B) du t-PA, et un activateur du plasminogène hybride à simple chaîne comprenant une chaîne A consistant essentiellement en le domaine de la boucle 2 du t-PA ou les domaines du doigt et de la boucle 2 du t-PA, liés en série à la région catalytique (chaîne B) de l'u-PA, où la jonction entre le(s) domaine(s) de chaîne A et la chaîne B est sur le site d'activation.
- 15 15. Composition pharmaceutique selon la revendication 11, comprenant en tant que composant A un activateur du plasminogène hybride à simple chaîne choisi parmi [uPA(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] et [tPA(1-3)-tPA(176-275)-uPA(159-411)].
- 20 16. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un mutant d'activateur du plasminogène hybride u-PA/t-PA, dans lequel au moins l'un des sites de N-glycosylation est modifié de telle sorte que la glycosylation ne puisse se produire sur ce site.
- 25 17. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant B une hirudine choisie parmi la variante d'hirudine HV1 ayant la formule

40 Val Val Tyr Thr Asp Cys Thr Glu Ser Gly
Gln Asn Leu Cys Leu Cys Glu Gly Ser Asn
Val Cys Gly Gln Gly Asn Lys Cys Ile Leu
Gly Ser Asp Gly Glu Lys Asn Gln Cys Val
45 Thr Gly Glu Gly Thr Pro Lys Pro Gln Ser
His Asn Asp Gly Asp Phe Glu Glu Ile Pro
W
Glu Glu Tyr Leu Gln

(III)

où W représente le groupe hydroxy phénolique de Tyr (désulfato hirudine HV1) ou un groupe de formule -O-SO₃H (hirudine HV1), la variante d'hirudine HV2 ayant la formule



Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 5 Ile Leu Gly Ser Asn Gly Lys Gly Asn Gln Cys Val Thr Gly
 Glu Gly Thr Pro Asn Pro Glu Ser His Asn Asn Gly Asp Phe
 Glu Glu Ile Pro Glu Glu Tyr Leu Gln

(IV)

et la variante d'hirudine PA ayant la formule

Ile Thr Tyr Thr Asp Cys Thr Glu Ser Gly Gln Asn Leu Cys
 Leu Cys Glu Gly Ser Asn Val Cys Gly Lys Gly Asn Lys Cys
 15 Ile Leu Gly Ser Gln Gly Lys Asp Asn Gln Cys Val Thr Gly
 Glu Gly Thr Pro Lys Pro Gln Ser His Asn Gln Gly Asp Phe
 Glu Pro Ile Pro Glu Asp Ala Tyr Asp Glu

(V)

ou un fragment ou un mutant de l'une quelconque des dites variantes d'hirudine conservant l'activité antithrombotique.

18. Composition pharmaceutique selon la revendication 17, comprenant en tant que composant B une hirudine choisie parmi la désulfato hirudine HV1, l'hirudine HV1, l'hirudine HV2 et l'hirudine PA.

19. Composition pharmaceutique selon la revendication 18, comprenant en tant que composant B la désulfato hirudine HV1.

20. Composition pharmaceutique selon la revendication 17, comprenant en tant que composant B une hirudine choisie parmi la [Ile^{1,2}]-désulfato hirudine HV1, la [Gly¹]-désulfato hirudine HV1, la [Gln^{57,58,61,62}]-désulfato hirudine HV1, la [Pro⁶⁶]-désulfato hirudine HV1, la [Gln²⁷,Arg⁴⁷]-désulfato hirudine HV1, la [Met⁰,Gln²⁷,Arg⁴⁷]-désulfato hirudine HV1, la [Gln²⁷,Gln³⁶,Arg⁴⁷]-désulfato hirudine HV1, la [des-Val¹,Thr²]-désulfato hirudine HV1 et la [Lys⁴⁷]-hirudine HV2.

21. Composition pharmaceutique selon la revendication 1, comprenant en tant que composant A un activateur du plasminogène choisi parmi scu-PA, t-PA, [Ala⁴⁵⁰]-t-PA, [Asn¹¹⁹, Ala¹⁸⁶, Ala⁴⁵⁰]-t-PA, [uPA-(1-44)-tPA(176-527)], [tPA(1-49)-tPA(176-275)-uPA(159-411)] et [tPA(1-3)-tPA(176-275)-uPA(159-411)] , et en tant que composant B de la désulfato hirudine HV1.

22. Composition pharmaceutique selon la revendication 1 pour utilisation dans une méthode de traitement du corps humain ou animal.

23. Composition pharmaceutique selon la revendication 1 pour utilisation dans la prévention ou le traitement de la thrombose ou d'affections provoquées par une thrombose, une artériosclérose, un infarctus du myocarde et un infarctus cérébral, une thrombose veineuse, une thromboembolie, une thrombose post-chirurgicale et une thrombophlébite.

24. Composition pharmaceutique selon la revendication 1 pour utilisation dans la prévention ou le traitement de l'infarctus du myocarde.



25. Composition pharmaceutique selon la revendication 1, dans laquelle le rapport en poids entre le composant A et le composant B est entre 20:1 et 2:1.

5 26. Composition pharmaceutique selon la revendication 25, dans laquelle le rapport en poids entre le composant A et le composant B est entre 5:1 et 2:1.

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